WIZZ

Page 1 02/08/99 M.BORIN 17 FILE USPATFULL 13 12 FILE WPIDS L14 162 FILE BIOSIS L15 83 FILE EMBASE L16 117 FILE MEDLINE L17 L18 137 FILE CAPLUS 65 FILE SCISEARCH L19 O FILE INVESTEXT L20 6 FILE DRUGU L21 TOTAL FOR ALL FILES 599 S LYSOSTAPHIN (10A) (ANTIMICROB#### OR MICROB### OR STAPHYLOCOC L22 280 DUPLICATE REMOVE L15-L19 (284 DUPLICATES REMOVED) L23 => d 114 bib, abs 1-14COPYRIGHT 1999 DERWENT INFORMATION LTD L14 ANSWER 1 OF 12 WPIDS AN 98-332208 [29] WPIDS 1020 DNC C98-102903 Treatment of staphylococcal infections, especially mastitis uses recombinant bacteriocin lysostaphin originally from Staphylococcus simulans. DC B04 B05 C03 C06 D16 BLACKBURN, P; POLAK, J TN (AMBI-N) AMBI INC PΑ CYC 1 US 5760026 A 980602 (9829)\* 10 pp US 5760026 A CIP of US 87-48412 870511, Cont of US 88-188183 880428, Cont PΙ of US 90-535286 900608, Cont of US 92-935121 920820, US 94-303551 940909 870511; US 90-535286 880428; US 87-48412 PRAI US 88-188183 920820; US 94-303551 940909 US 92-935121 WPIDS 98-332208 [29] AN UPAB: 980722 US 5760026 A AB Treatment of recurring staphylococcal mastitis resulting from intracellular Staphylococcus aureus comprises administering to an infected gland, by intramammary infusion, a therapeutic agent consisting of bacteriocin lysostaphin produced by recombinant means, in a pharmaceutical carrier in an amount sufficient to eliminate recurring staphylococcal mastitis. USE - The method is useful for the treatment of staphylococcal infections, especially bovine mastitis, by the administration of the bacteriocin lysostaphin. Mastitis is caused by infection of bovine milk glands by S.aureus and S.agalactiae (and some Gram negative bacteria, to a lesser extent), leading to decreased and unusable milk production and in extreme cases, death. The bacteriocin lysostaphin is a protein that is produced by S.simulans (NRRL B-2628) that kills and lyses related bacteria. Antibacterial compositions can be applied to teats for prophylactic use. ADVANTAGE - Previous mastitis treatments rely on antibiotics where

the disease has we poor response to treatment, even re-occurring. Extensive treatment this way can also lead to antistrains of Staphylococcus. Dwq.0/1COPYRIGHT 1999 DERWENT INFORMATION LTD ANSWER 2 OF 12 WPIDS L1496-117492 [13] WPIDS DNC C96-037290 New lysostaphin gene with deletion in pro-segment repeat region - for prodn. of Lactobacillus strains useful in prodn. of fermented foods and as protective cultures to inhibit growth of Staphylococcus. DC D13 D16 CAVADINI, C; HAMMES, W; HERTEL, C (MUEL-N) MUELLER & CO KG KARL; (MUEL-N) MUELLER GMBH & CO KARL CYC 12 DE 4425645 A1 960222 (9613)\* 19 pp A1 970226 (9714)# DE EP 759473 R: AT CH DK ES FR GB IE IT LI NL SE ADT DE 4425645 A1 DE 94-4425645 940720; EP 759473 A1 EP 95-113211 950823 PRAI DE 94-4425645 940720; EP 95-113211 WPIDS 96-117492 [13] UPAB: 960329 DE 4425645 A New lysostaphin gene (A) of Staphylococcus simulans has a deletion in the repeat region of the pro segment. Also new are plasmids and microorganisms contg (A). USE - The modified microorganisms are useful in fermentation (e.g. prodn. of meat prods.) and as protective cultures, against contamination by Staphylococcus in foods, (e.g. mayonnaise or milk products). ADVANTAGE - Modified (A) can be expressed in food grade microorganisms, i.e. they produce active lysostaphin which is able to lyse most Staphylococci including food contaminants such as S. aureus. (A) expresses a protein with the normal pre-sequence, a truncated pro-sequence and a mature lysostaphin sequence. In Lactobacillus the signal and residual pro sequences are removed. The new cells do not integrate (A) into the genome and since they gradually lose the (A)-contg. plasmid can be released into the environment without danger. Dwg.0/10 COPYRIGHT 1999 DERWENT INFORMATION LTD L14 ANSWER 3 OF 12 WPIDS 94-148921 [18] WPIDS ANDNC C94-068635 Prepn. of restricting endonuclease SAU 6782 - by culturing microorganism Staphylococcus aureus 6782. treating with lysostaphin, ultrasonically disintegrating and removing nucleic acids. DC ARUTYUNYAN, E E; GONCHAR, N A; NIKOLSKAYA-SANOVICH, I I IN (BIOL-R) BIOLOG MED CHEM INST; (VACC-R) VACCINE & SERUM INST PΑ CYC 1 SU 1796676 A1 930223 (9418)\* 5 pp SU 1796676 A1 SU 91-4920752 910320 PRAI SU 91-4920752 910320 AN 94-148921 [18] WPIDS SU 1796676 A UPAB: 940622 Method involves culturing the microorganism Staphylococcus aureus 6782, treating the bacterial mass with lysostaphin, ultrasonic disintegration, removal of the nucleic acids by streptomycin sulphate, salting out of the proteins by ammonium sulphate, and cation exchange chromatography on R11 phosphocellulose. The cation exchange chromatography is carried out directly after salting out of the proteins, using a double combined increasing NaCl concn. gradient from 0.0 to 1.0 M, and pH values decreasing from 9.0 to

USE/ADVANTAGE - In biotechnology and genetic engineering, and in

6.0.

molecular biology 1, genetic experiments in studying the structure and functions of DNA and the construction of recombilat molecules.

In an example, cells of Staphylococcus aureus 6782 were cultured in a

medium based on a casein trypsin hydrolysate and yeast water, and deposited by centrifuging. The bacterial mass was suspended in the working buffer (pH 7.4) contg. potassium phosphate, beta-mercaptoethanol, and EDTA, lysostaphin was added, and the mass was incubated at 20 deg.C. The cell suspension obtd. was subjected to ultrasonic disintegration and centrifuged at 105 thousand g for 1 hr, to give a supernatant liquid in the form of a crude extract, from which the nucleic acids were removed by means of streptomycin sulphate. The proteins were then salted out with ammonium sulphate, the deposit was dissolve din the buffer, the ammonium sulphate was dialysed out, and the soln. was subjected to column chromatography, with an NaCl concn. gradient increasing from 0.0 to 1.0 M, and the pH decreasing from 9 to 6. Restrictase was eluted in a narrow peak in the region of 0.4-0.45 M NaCl, and non-specific nucleases at 0.45-0.55 NaCl concn. The R Sau 6782 preparation obtained contained an insignificant quantity of non-specific impurities, and the enzyme yield was 1300 units from 1 g of biomass, specific enzyme activity 60000 units per mg. Dwg.0/0

L14 ANSWER 4 OF 12 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD

AN 94-011014 [02] WPIDS

DNC C94-004489

Purificn. of lysostaphin endopeptidase - comprises absorbing LEP on microbe body having dissolution resistance against LEP.

DC B04 D16

PA (SAOC) MERCIAN CORP

CYC 1

PI JP 05317045 A 931203 (9402)\* 4 pp

ADT JP 05317045 A JP 92-148955 920518

PRAI JP 92-148955 920518

AN 94-011014 [02] WPIDS

AB JP05317045 A UPAB: 940223

In the method, the LEP is adsorbed on a microbe body of a microbe having dissolution resistance against LEP or a treated prod. and then eluted. The microbe is pref. of Staphylococcus genus.

USE/ADVANTAGE - The method can prepare a prod. of high purity in a high yield very easily.

In an example, Staphylococcus aureus Kowa I (ATCC12598) was cultured in 50ml of a brain-heart infusion broth at 37 deg.C for 24 hours. The culture was centrifuged and the microbe body was suspended in PBS buffer to 108 cells/ml. LEP was added to it to 4 U/ml and cultured at 37 deg.C overnight. It was sepd. to a single colony in a nutrient agar medium to give a colony of LEP-resistant microbe, Staphylococcus aureus MY1. It was inoculated to 50ml of a tryptic soy broth and cultured at 37 deg.C for 24 hours. The culture was centrifuged and the microbe body was washed with 500 ul 50 mM Tris-HCl buffer and suspended in 100 ml 3M K thiocyanate and heated at 100 deg.C for 30 min.. The microbe body was collected by centrifugation and washed with 100 ml 3Mk thiocyanate and then with 500 ml 50 mM Tris-HCl buffer and then suspended in 100 ml of the same buffer. The suspension was added to a culture supernatant contg. LEP to  $1\ x\ 10$  power 9cells/ml and the mixture was stirred at 4 deg.C for 1 hour to absorb LEP specifically. LEP was eluted from it and the LEP soln. was dialysed and freeze-dried to give 7330 U of highly pure LEP. Dwg.0/0

L14 ANSWER 5 OF 12 WPIDS COPYRIGHT 1999 DERWENT INFORMATION LTD

AN 93-054352 [07] WPIDS

DNC C93-024311

TI Detection of methicillin-resistant staphylococci - using polymerase chain reaction method, and DNA primers, for rapid, sensitive and accurate detection.

```
DC
     B04 D16
     SKATRUD, P L; UN
IN
     (ELIL) LILLY & CO ELI
PA
CYC 18
               A1 930217 (9307)* EN
                                        16 pp
    EP 527628
PΙ
     CA 2075423 A 930214 (9318)
     JP 05329000 A 931214 (9403)
                                        12 pp
                                        17 pp
     EP 527628 B1 960703 (9631)
                                  EN
         R: AT BE CH DE DK ES FR GB GR IE IT LI LU NL PT SE
     DE 69211921 E 960808 (9637)
     ES 2089409 T3 961001 (9645)
ADT EP 527628 A1 EP 92-307307 920810; CA 2075423 A CA 92-2075423 920806; JP
     05329000 A JP 92-214968 920812; EP 527628 B1 EP 92-307307 920810; DE
     69211921 E DE 92-611921 920810, EP 92-307307 920810; ES 2089409 T3 EP
     92-307307 920810
FDT DE 69211921 E Based on EP 527628; ES 2089409 T3 Based on EP 527628
                    910813
PRAI US 91-744770
                      WPIDS
     93-054352 [07]
ΑN
                    UPAB: 931119
     EP 527628 A
AΒ
     Method comprises (a) performing polymerase chain reaction (PCR) on
     samples, the PCR being primed by DNA primers, the DNA primers being
     composed of 2 oligonucleotides of high GC content, where one
     oligonucleotide has DNA sequence comprised of the coding strand of
     Staphylococcus mecA gene and the second DNA primer has DNA sequence
     comprised of the non-coding strand of a Staphylococcus mecA gene and (b)
     analysing the reaction prod.
          Also claimed are (i) method for the rapid release of DNA from
     Staphylococci comprising (a) treating a sample contg.
     Staphylococci with lysostaphin, (b) treating the
     resulting sample with proteinase K and (c) incubating the resulting sample
     in a boiling water bath, (ii) method for detecting methicillin resistant
     staphylococcal infections in sample, comprising (a) carrying out steps
     (a)-(c) and (b) carrying out steps (a)-(b).
          ADVANTAGE - The method can be used for the rapid, sensitive and
     accurate detection of methicillin-resistant Staphylococcal infections
     caused by e.g. methicillin-resistant S. aureus (MRSA) or
     methicillin-resistant S. epidermis (MRSE)
     Dwg.0/0
                    UPAB: 960808
ABEQ EP 527628 B
     A method for detecting methicillin resistant staphylococcal infections,
     said method comprising: a) performing the polymerase chain reaction on
     clinical samples suspected of staphylococcal infection, said polymerase
     chain reaction being primed by DNA primers, said DNA primers being
     composed of two oligonucleotides of high GC content, wherein one
     oligonucleotide has a DNA sequence comprised by the coding strand of a
     Staphylococcus mecA gene and the second DNA primer has a DNA sequence
     comprised by the non-coding strand of a Staphylococcus mecA gene; and b)
     analyzing the reaction product of step a.
     Dwg.0/0
                             COPYRIGHT 1999 DERWENT INFORMATION LTD
L14 ANSWER 6 OF 12 WPIDS
     92-151734 [19]
                      WPIDS
AN
DNC C92-070227
     Lysostaphin prodn. - by culture of Staph. simulans on medium contg.
     pancreatic casein peptone, fractional pptn., dialysis and chromatography.
DC
     B04 D16
     PAUL, P; REISSBRODT, R
IN
     (REIS-I) REISSBRODT R
PΑ
CYC 1
     DE 4033752 A 920430 (9219)*
                                          3 pp
ΡI
ADT DE 4033752 A DE 90-4033752 901024
 PRAI DE 90-4033752 901024
```

WPIDS

92-151734 [19]

AN

```
: 931006
                     in (I), a staphylococci-lysing
   enzyme complex from Staph. simulans var. staphylolficus comprises (1)
  DE 4033752 A
   adding pancreatic casein peptone (A), chosen according to its analysed
   nutrient parameters, to a standard nutrient medium for growing the prodn.
   strain so as to regulate the subsequent purification process, (2)
   pre-purification of (I) by fractional (NH4)SO4 pptn. from the culture
   supernatant, (3) removing DNase from crude (I) by treatment with urea and
   dialysis, (4) stabilising the labile (I) complex (isolated by
   chromatography using a IMNaCl gradient) with protective colloids (e.g.
        USE/ADVANTAGE - (I) is used for simple and rapid differentiation
    gelatine-sucrose) and freeze drying.
    between staphylococci (lysed) and micrococci (resistant) and can also be
    used in bacterial-genetic investigation and for isolation of bacterial
    metabolite. The method provides (I) of high activity and free of DNase.
    (0/3)
                            COPYRIGHT 1999 DERWENT INFORMATION LTD
    0/3
L14 ANSWER 7 OF 12 WPIDS
    Novel lanthionine contg. bacteriocin and lysostaphin compsns. - useful as
NA
DNC C90-125224
     enhanced broad range bactericides.
TI
     BLACKBURN, P; GUSIK, S; POLAK, J; RUBINO, S D; GUSIK, S A
     (PUBL-N) PUBLIC HEALTH RES INST NEW YORK; (MICR-N) APPLIED MICROBIOLOGY
DC
IN
     INC; (PUBL-N) PUBLIC HEALTH RES
PΑ
     WO 9009739 A 900907 (9038)*
 CYC 16
      AU 9052850 A 900926 (9050)
 ΡI
      ZA 9001499 A 901128 (9102)
US 4980163 A 901225 (9103)
                    901031 (9107)
      FI 9005378 A
      NO 9004729 A 901115 (9114)
      EP 424484 A 910502 (9118)
                     910628 (9131)
                  Т
      ни 55607
      CS 9000984 A 910716 (9143)
      JP 03504864 W 911024 (9149)
                 A 921028 (9301)
      NZ 232700
                  A9 940630 (9431)
       DD 301903
                  B1 940810 (9431)
                                    ΕN
                                          10 pp
       EP 424484
       DE 69011460 E 940915 (9436)
                 A 950315 (9517)
       IL 93527
                 B6 950315 (9520)
       CZ 279273
                 B 950823 (9542)
       IE 64710
       RU 2048151 C1 951120 (9629)
       ZA 9001499 A ZA 90-1499 900227; US 4980163 A US 89-317627 890301; EP
        424484 A EP 90-904988 900227; JP 03504864 W JP 90-504798 900227; NZ 232700
        A NZ 90-232700 900227; DD 301903 A9 DD 90-338282 900301; EP 424484 B1 EP
        90-904988 900227, WO 90-US1053 900227; DE 69011460 E DE 90-611460 900227,
   ADT
        EP 90-904988 900227, WO 90-US1053 900227; IL 93527 A IL 90-93527 900226;
        CZ 279273 B6 CS 90-984 900301; IE 64710 B IE 90-721 900228; RU 2048151 C1
        SU 90-4831853 900227; CA 2028140 C CA 90-2028140 900227
   FDT EP 424484 B1 Based on WO 9009739; DE 69011460 E Based on EP 424484, Based
        on WO 9009739; CZ 279273 B6 Previous Publ. CS 9000984
                        890301
    PRAI US 89-317627
         Novel compsn. comprising lysostaphin and a lanthionine contg. bacteriocin.
    NΑ
              Pref. lanthionic bacteriocin is selected from rosin,
    AΒ
         New broad range antibacterial compsn. comprises lysostaphin and
    ABEQ US 4980163 A UPAB: 930928
```

lanthionine control bacteriocin opt. with chelator and surfactant. Pref. bacter cin is nisin, subtilin, epider 1, cinnamycin, duramycin, ancovenin or Pep 5, and chelators include EDTA, with triton or Tween etc. as surfactant. USE/ADVANTAGE - More effective bactericidall effect against broad range microbial infections esp. Gram +- than lysostaphin alone. Effective conc. 0.1-100mgc/ml lysostaphin: 0.1-300 mcg/ml nisin; 0.1-20 mM chelator; 0.01-1 % vol. surfactant. @ UPAB: 940921 ABEQ EP 424484 B A composition comprising lysostaphin and a lanthionine containing bacteriocin. Dwg.0/0 COPYRIGHT 1999 DERWENT INFORMATION LTD ANSWER 8 OF 12 WPIDS WPIDS ΑN 89-324396 [44] C89-143659 DNC Compsn. for preventing staphylococcal infections - contains TIlysostaphin, surfactant and penicillin for synergistic effect. B05 C03 D16 DC BLACKBURN, P; POLLAK, J; POLLACK, J IN (MICR-N) APPLIED MICROBIOLOGY INC; (PUBL-N) PUBLIC HEALTH RES PΑ CYC ZA 8806446 A 890830 (8944)\* 38 pp PΙ EP 359873 ) A 900328 (9013)# EN R: AT BE CH DE ES FR GB GR IT LI LU NL SE A 900330 (9017)# PT 88472 JP 02083336 A 900323 (9018)# AU 8821793 A 900308 (9019)# DK 8804846 A 900301 (9019)# 910528 (9127)# HU 55229  $\mathbf{T}$ EP 359873 B1 930915 (9337)# EN R: AT BE CH DE FR GB IT LI LU NL SE DE 3884200 G 931021 (9343)# JP 06045553 B2 940615 (9422)# 13 pp CA 1330758 C 940719 (9434)# B 950322 (9521)# IE 63009 A 951231 (9614)# IL 87686 ZA 8806446 A ZA 88-6446 880830; EP 359873 A EP 88-308667 880919; JP 02083336 A JP 88-234685 880919; EP 359873 B1 EP 88-308667 880919; DE 3884200 G DE 88-3884200 880919, EP 88-308667 880919; JP 06045553 B2 JP 88-234685 880919; CA 1330758 C CA 88-575957 880829; IE 63009 B IE 88-2675 880905; IL 87686 A IL 88-87686 880906 FDT DE 3884200 G Based on EP 359873; JP 06045553 B2 Based on JP 02083336 880830; EP 88-308667 880919; JP 88-234685 880919; PRAI ZA 88-6446 880829; IE 88-2675 880905; DE 88-3884200 880919; CA 88-575957 IL 88-87686 880906 ΑN 89-324396 [44] WPIDS UPAB: 931129 ZA 8806446 A AΒ Compsn. (A) for killing staphylococci comprises lysostaphin (I) and synergist(s) from penicillin, synthetic penicillins, other antibiotics, chelating agents, mild surfactants and other membrane active agents, in amts. effective to kill staphylococci. Preventing bovine mastitis comprises dipping teats in a soln. (B) of 0.01-10.0 mcg/ml of (I) in a suitable carrier, before and after each milking. The compsn. (A) and teat dip soln. (B) may also comprise lysozyme and mutanolysin. USE/ADVANTAGE - Used for treatment and prevention of staphylococcal infections esp. bovine mastitis. (I) is effective even against chronic mastitis without adverse immunogenic effects. The compsn. is highly synergistic, with potentiation of (I) by e.g. 1000 times or more when

surfactants are added. The compsn. may also be infused into the infected udder, and is also useful in wound dressings and medications, disinfectant scrubs, wipes or lotions, and in surgical implants. Compsns. may also be

```
used for cleaning edical instruments, surfaces, adding, etc., related uses (treament of meat, eggs, cheese and lish or food packaging
     and handling appts.), and for nasal infusion to reduce intranasal carriage
     of Staphylococci. (Provisional Basic previously advised in week 8939)
     Dwg.0/1
                    UPAB: 931123
ABEQ EP 359873 B
     A lysostaphin-containing composition for killing
     staphylococci (and which therefore does not itself contain
     staphylococci), characterised in that it also comprises at least one cell
     wall-active antiobiotic in an amount effective synergistically to enhance
     the bactericidal effect of lysostaphin against
     staphylococcal mastitis.
     Dwq.0/1
                             COPYRIGHT 1999 DERWENT INFORMATION LTD
L14 ANSWER 9 OF 12 WPIDS
                      WPIDS
     89-216318 [30]
ΑN
DNC C89-096165
     Genetic DNA - used for prodn. of protein which binds with Fc part of
     immunoglobulin G.
DC
     B04 D16
     (SAOC) SANRAKU OCEAN CO LTD
CYC 1
     JP 01153093 A 890615 (8930)*
                                          7 pp
ADT JP 01153093 A JP 87-311037 871210
                    871210
PRAI JP 87-311037
     89-216318 [30]
                      WPIDS
AN
     JP01153093 A UPAB: 930923
AB
     Genetic DNA is claimed which relates to prodn. of protein that can bind
     with the Fc part of immunoglobulin G. The scale of the molecule is ca.
     1.05 kb, and it has a sensitivity against restriction enzymes, namely (a)
     has one recognition site by ClaI, ScaI, SalI, Sau3A, EcoRI, SacI, AccI,
     and (b) is not cleaved by BglII, EcoRV, STuI, PvUII, HpaI, XhoI, BclI,
     KpnI, Sph.
          Pref. the protein A-like substance is SRP-2 (m.w.: 24,000, i.p.: pH
     4.5, U.V. 275 nm max. (E 1%/1 cm = 1.60); binding affinity with human IgG
     11 mg/mg). Cells of Staphylococcus aureus SR-1 are lysed with
     lysostaphin, and whole DNA is extracted. It is partly decomposed
     with Sau3A, and 2-10 kb DNA is purified and obtd. by agarose
     electrophoresis. Plasmid vector pUC8 is decomposed with BamHI, and both
     decomposed fragments are combined and ligated with T4 ligase. Thus obtd.
     recombinant plasmid is used for transformation of E.coli JM103, then a
     plasmid reserving colony is selected by ampicillin contg. selective
     medium, (E. coli 7-3, FERM-9746).
          USE/ADVANTAGE - By transforming various host bacteria with the
     genetic RNA contg. recombinant plasmid, protein A-like protein
     producibility can be expressed to the transformant, and is useful for
     prodn. of the protein.
L14 ANSWER 10 OF 12 WPIDS
                               COPYRIGHT 1999 DERWENT INFORMATION LTD
     87-306856 [43]
                       WPIDS
ΑN
     C87-130725
     Recombinant plasmids contg. gene for lysostaphin - a bactericide
     specific for staphylococci, and transformed microbial
     hosts.
     B04 D16
DC
     RECSEI, P A
IN
      (PUBL-N) PUBLIC HEALTH RES; (MICR-N) APPLIED MICROBIOLOGY INC; (PUBL-N)
     PUBLIC HEALTH RES INST INC CITY NEW YORK
CYC 21
     WO 8706264 A 871022 (8743) * EN
PΙ
         RW: AT BE CH DE FR GB IT LU NL SE
```

W: AU DK FI HU JP ZA 8702687 A 871116 (8804)

```
79 (8805)
    AU 8773021 A
                        21 (8822)
                A 8
    PT 84705
    DK 8706592 A 871215 (8834)
               A 890125 (8904)
                                  EN
     EP 299978
         R: AT BE CH DE FR GB IT LI LU NL SE
     FI 8804691 A 881012 (8927)
     JP 01503036 W 891019 (8948)
              T 900328 (9019)
    HU 50874
    US 4931390 A 900605 (9026)
     CA 1297051 C 920310 (9216)
              A 920818 (9244)
     IL 82260
     EP 299978 B1 930804 (9331) EN
                                       22 pp
        R: AT BE CH DE FR GB IT LI LU NL SE
     DE 3786918 G 930909 (9337)
                B 960229 (9613)
     FI 96322
    WO 8706264 A WO 87-US873 870415; ZA 8702687 A ZA 87-2687 870414; EP 299978
     A EP 87-903128 870415; JP 01503036 W JP 87-502621 870415; US 4931390 A US
     87-34464 870410; IL 82260 A IL 87-82260 870415; EP 299978 B1 EP 87-903128
     870415, WO 87-US873 870415; DE 3786918 G DE 87-3786918 870415, EP
     87-903128 870415, WO 87-US873 870415; FI 96322 B WO 87-US873 870415, FI
     88-4691 881012
    EP 299978 B1 Based on WO 8706264; DE 3786918 G Based on EP 299978, Based
     on WO 8706264
                    870410; US 86-852407
                                          860416
PRAI US 87-34464
     87-306856 [43]
                     WPIDS
ΑN
     WO 8706264 A
                   UPAB: 960520
ÆΒ
     Recombinant plasmids able to express a gene coding for lysostaphin
     (I) in a transformed microbial host are new. Also new are (1)
     the transformed microorganisms; (2) pure (I) free of immunogenic
     staphylococcal contaminants, and (3) a 1.5 kb DNA fragment coding for (I).
          Hosts are E.coli K-12 JM105 (transformed with plasmid pRG5); Bacillus
     sphaericus strain 00 (plasmid pJP1) and B.subtilis DB170 (plasmids, pJP1,
     pDF8 or pRP1). The 1.5 kb fragment (complete sequence reproduced)
     comprises promoter sequences (nucleotides 89-95 and 110-119);
     ribosome-binding site (231-235) and an open reading frame from TTG
     (245-247) to TGA (1412-1414), coding for preprolysostaphin (PPL). PPL is a
     precursor for mature, active (I) and contains a signal peptide and
     prelyostaphin (PL) which is processed to mature (I) by cleavage of the
     Arg-Ala (143-144) bond in PL. PPL contains 389 amino acids.
          USE/ADVANTAGE - (I) is a bacteriocin which kills most Staphyloccal
     species but not other bacteria. The transformants produce (I) identical
     with the natural material and sometimes at a much higher level than S.
     simulans.
     0/1
     Dwg.0/1
                    UPAB: 930922
ABEQ US 4931390 A
     Recombinant plasmids contain a DNA sequence which codes for lysostaphin,
     which expresses a gene encoding lysostaphin from staphyloccus
     simulens (NRRL 2-2228) in transformant microbial hosts. Pref.
     recombinant plasmid comprises pRGS, pJP1, pDF8, or pRP1. Transformed
     microorganism comprises E.coli a yeast, Streptomyces, spp., or Bacillus
     ssp.
          ADVANTAGE - Bacillus sphaericus strain oo/pJp1 transformants produce
     5 times the amt. of prod. as S.simulans.
ABEQ EP 299978 B
                  UPAB: 931118
     A 1.5 kilobase DNA coding for lysostaphin, which comprises a nucleotide
     sequence of Formula I: having an open reading frame extending from a TTG
     initiation codon at nucleotides 245-247 to a TGA termination codon at
     nucleotides 1412-1414, the open reading frame coding for
     prepropysostaphin, which is a precursor to mature active lysostaphin,
     having a sequence of 389 amino acids which comprises a signal peptide and
```

prolysostaphin, which is processed after synthesis to mature lysostaphin.

Dwg.0/1

```
COPYRIGHT 1999 DERWENT 1
L14 ANSWER 11 OF 12
                        IDS
                     WPIDS
     72-67951T [43]
     Lysostaphin - from cultivation of staphylococcus staph
     ylolyticus in media contg enzymatically hydrolysed casein.
DC
     (BRIM) BRISTOL-MYERS CANADA LTD
PA
CYC
                           (7243)*
PΙ
     CA 911917
               Α
                    680517
PRAI CA 68-20358
     72-67951T [43] WPIDS
     CA 911917 A UPAB: 930831
AΒ
     Lysostaphin is produced by cultivating a strain of
     Staphylococcus staphylolyticus in an aq. nutrient medium contg. C
     and N supplying nutrients. The medium contains >=4, pref. 4-10 % wt
     enzymatically hydrolysed casein as N-supplying nutrient and its pH is
     6.5-8.5. The C source is pref. >=0.5% wt glycerol, mannose or galactose.
     The product is an antibiotic which specif. lyses Staphylococcus organisms.
     By this method an increased yield of product is possible and the need for
     pH adjustment is eliminated if the prefd. C source is used.
                              COPYRIGHT 1999 DERWENT INFORMATION LTD
L14 ANSWER 12 OF 12 WPIDS
                      WPIDS
AN
     66-33531F [00]
     Fermentative production of lysostaphin antibiotic.
ΤI
DC
     (MEAD) MEAD JOHNSON & CO
PA
CYC 8
                           (6800)*
     US 3398056 A
PI
                           (6801)
     BE 716314
                Α
     FR 1574204 A
                           (6801)
     NL 6807706 A
                           (6801)
     GB 1196835 A
                           (7026)
                           (7137)
     CH 509408
                Α
     JP 47013719 B
                           (7217)
     DE 1767687 A 710902 (8517)
PRAI US 64-381684
                    640710
     66-33531F [00]
                      WPIDS
ΑN
     US 3398056 A
                    UPAB: 930831
AΒ
     A process for the production of lysostaphin by the fermentation
     of Staphylococcus staphylolyticus in an aqueous nutrient medium,
     in which one or both of the following improvements are
     incorporated in the process:
           (1) the fermentation is carried out at pH6.5-8.5 in the
     presence of at least 4% w/w enzymatically hydrolysed casein.
           (2) the fermentation is carried out at pH 6.4-8.5 in the
     presence of at least 0.5% of glycerol, mannose or galactose.
           Antibiotic.
```

```
17 FILE USPATFULL
L13
            12 FILE WPIDS
L14
           162 FILE BIOSIS
L15
            83 FILE EMBASE
L16
            117 FILE MEDLINE
L17
            137 FILE CAPLUS
L18
             65 FILE SCISEARCH
L19
              O FILE INVESTEXT
L20
              6 FILE DRUGU
L21
     TOTAL FOR ALL FILES
            599 S LYSOSTAPHIN (10A) (ANTIMICROB#### OR MICROB### OR STAPHYLOCOC
L22
            280 DUPLICATE REMOVE L15-L19 (284 DUPLICATES REMOVED)
L23
```

=> d 123 an, ti 50-280

- L23 ANSWER 50 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- 1992:520872 BIOSIS AN
- INHIBITORY EFFECT OF 5 QUINOLONES ON DNA GYRASE FROM STAPHYLOCOCCUS-ΤI AUREUS.
- L23 ANSWER 51 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- 1993:204544 BIOSIS ΑN
- Sensitivity of Dermatophilus congolensis to antibiotic substances of ΤI staphylococci.
- L23 ANSWER 52 OF 280 CAPLUS COPYRIGHT 1999 ACS
- 1992:229239 CAPLUS ΑN
- 116:229239 DN
- Cloning and expression of the lysostaphin gene in Bacillus subtilis and Lactobacillus casei
- L23 ANSWER 53 OF 280 CAPLUS COPYRIGHT 1999 ACS
- 1992:124351 CAPLUS AN
- 116:124351 DN
- EGTA inhibition of DNase activity in commercial lysostaphin preparations ΤI
- L23 ANSWER 54 OF 280 CAPLUS COPYRIGHT 1999 ACS
- ΑN 1992:231723 CAPLUS
- 116:231723 DN
- Purification and application of lysostaphin
- L23 ANSWER 55 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS . DUPLICATE 29
- 1992:93289 BIOSIS ΑN
- LYSOSTAPHIN USE OF A RECOMBINANT BACTERICIDAL ENZYME AS A MASTITIS TΙ THERAPEUTIC.
- L23 ANSWER 56 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1991:486327 CAPLUS

- L23 ANSWER 57 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1993:3667 CAPLUS
- DN 118:3667
- TI Biochemical and genomic characteristics of Micrococcaceae from French dry sausages
- L23 ANSWER 58 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 30
- AN 1991:459340 BIOSIS
- VISUALIZATION OF ENDO-BETA-N-ACETYLGLUCOSAMINIDASE LYSOZYME AND LYSOSTAPHIN AFTER POLYACRYLAMIDE GEL ELECTROPHORESIS IN THE PRESENCE OF SDS.
- L23 ANSWER 59 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.DUPLICATE 31
- AN 91012053 EMBASE
- TI .beta.-Lactamase is encoded on plasmid pACK3 in Staphylococcus simulans biovar staphylolyticus.
- L23 ANSWER 60 OF 280 MEDLINE
- AN 91169234 MEDLINE
- TI Beta-lactamase is encoded on plasmid pACK3 in Staphylococcus simulans biovar staphylolyticus.
- L23 ANSWER 61 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 32
- AN 1991:431094 BIOSIS
- TI SEQUENCE ANALYSIS OF A STAPHYLOCOCCUS-AUREUS GENE ENCODING A PEPTIDOGLYCAN HYDROLASE ACTIVITY.
- L23 ANSWER 62 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1992:55305 CAPLUS
- DN 116:55305
- TI Characteristics of extracellular protein production by a plasmidless derivative of Staphylococcus simulans biovar staphylolyticus
- L23 ANSWER 63 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1991:651266 CAPLUS
- DN 115:251266
- TI Heat stable nuclease contamination of lysostraphin (final report)
- L23 ANSWER 64 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1991:120163 CAPLUS
- DN 114:120163 -
- TI Lysostaphin digestion in preparation of vaccines
- L23 ANSWER 65 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1991:214519 CAPLUS
- DN 114:214519
- TI Synergistic bactericidal compositions comprising lysostaphin and a lanthionine-containing bacteriocin
- L23 ANSWER 66 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 33
- AN 1991:5783 BIOSIS
- TI EFFECT OF BITEK AGAR ON LYSOSTAPHIN SUSCEPTIBILITY OF STAPHYLOCOCCI.
- L23 ANSWER 67 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1990:526103 CAPLUS
- DN 113:126103

- TI The antibacterial activity of benzylpenicillin against Staphyrococcus aureus ingested granulocytes
- L23 ANSWER 68 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1990:492688 CAPLUS
- DN 113:92688
- TI New mold starter cultures by genetic modification
- L23 ANSWER 69 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1990:629004 CAPLUS
- DN 113:229004
- TI Solubilization of group- and type-specific streptococcal antigens with a murolytic enzyme from Staphylococcus hyicus
- L23 ANSWER 70 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 34
- AN 1991:28241 BIOSIS
- TI RAPID PURIFICATION METHOD OF LYSOSTAPHIN FOR ANALYSIS OF CELL WALL PROTEINS.
- L23 ANSWER 71 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 35
- AN 1990:423362 BIOSIS
- TI EFFECTS OF LYSOSTAPHIN ON STAPHYLOCOCCUS-AUREUS INFECTIONS OF THE MOUSE MAMMARY GLAND.
- L23 ANSWER 72 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1990:455476 CAPLUS
- DN 113:55476
- TI Biochemical characteristics of Staphylococcus species of human and bovine origin
- L23 ANSWER 73 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1990:494653 CAPLUS
- DN 113:94653
- TI Inhibition of the bacteriolytic effect of .beta.-lactam-antibiotics on Staphylococcus aureus by the polyanionic drugs suramin and Evans Blue
- L23 ANSWER 74 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1990:484861 CAPLUS
- DN 113:84861
- TI Bactericidal compositions containing lysostaphin
- L23 ANSWER 75 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1990:135576 CAPLUS
- DN 112:135576
- TI Process for the selective cleavage of fusion proteins with polyglycine-specific endoproteinase
- L23 ANSWER 76 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1989:630563 CAPLUS
- DN 111:230563
- TI Process for obtaining staphylococcal capsule polyosides and their use as vaccines and in diagnosis
- L23 ANSWER 77 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 36
- AN 1989:264380 BIOSIS
- TI SUSCEPTIBILITY OF METHICILLIN-RESISTANT STAPHYLOCOCCUS-AUREUS TO LYSOSTAPHIN.
- L23 ANSWER 78 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 37
- AN 1989:334426 BIOSIS
- TI PLASMID-ENCODED LYSOSTAPHIN ENDOPEPTIDASE RESISTANCE OF STAPHYLOCOCCUS-SIMULANS BIOVAR STAPHYLOLYTICUS.





1989:136303 BIO

ANTIBODY RESPONSE TO STAPHYLOCOCCUS-AUREUS SURFACE PROTEINS IN RABBITS WITH PERSISTENT OSTEOMYELITIS AFTER TREATMENT WITH DEMINERALIZED BONE IMPLANTS.

L23 ANSWER 80 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 39

1989:473586 BIOSIS

A COMPARISON OF THE RAPID THERMONUCLEASE TEST AND THE LYSOSTAPHIN SUSCEPTIBILITY TEST IN THE PRESUMPTIVE IDENTIFICATION OF STAPHYLOCOCCUS-AUREUS FROM POSITIVE BACTEC BLOOD CULTURES.

L23 ANSWER 81 OF 280 CAPLUS COPYRIGHT 1999 ACS

1989:492596 CAPLUS

111:92596

The spectrophotometric assay and kinetic properties of lysostaphin TI

L23 ANSWER 82 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS

1989:398226 BIOSIS

RESISTANCE OF STAPHYLOCOCCUS-SIMULANS BIOVAR STAPHYLOLYTICUS TO LYSOSTAPHIN ENDOPEPTIDASE IS PLASMID ENCODED.

L23 ANSWER 83 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS

1989:411960 BIOSIS

A LYSOZYME ISOLATED FROM RAINBOW TROUT ACTS ON MASTITIS PATHOGENS.

L23 ANSWER 84 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS

1989:335440 BIOSIS AN

USE OF STAPHYLOCOCCUS-AUREUS AND LYSOSTAPHIN IN AN ΤI ASSAY OF PHAGOCYTOSIS AND INTRACELLULAR KILLING.

L23 ANSWER 85 OF 280 CAPLUS COPYRIGHT 1999 ACS

1989:529497 CAPLUS AN

111:129497 DN

Prolysostaphin-processing protease from Staphylococcus TIsimulans--purification and some properties

L23 ANSWER 86 OF 280 CAPLUS COPYRIGHT 1999 ACS

1990:93181 CAPLUS ΝA

DN 112:93181

Cloning and expression of lysostaphin gene in Escherichia coli and TΙ Bacillus subtilis

L23 ANSWER 87 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS

1990:188813 BIOSIS

EXPRESSION OF LYSOSTAPHIN IN THE MILK OF TRANSGENIC ANIMALS TO COMBAT STAPHYLOCOCCAL MASTITIS.

L23 ANSWER 88 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 40

1988:436975 BIOSIS AN

DNA GYRASE OF STAPHYLOCOCCUS-AUREUS AND INHIBITORY EFFECT OF QUINOLONES ON ΤI ITS ACTIVITY.

L23 ANSWER 89 OF 280 CAPLUS COPYRIGHT 1999 ACS

1989:495036 CAPLUS AN

111:95036 DN

Removal of surface adherent Staphylococcus aureus in the determination of TΙ phagocytosis and intracellular killing by the use of lysostaphin

L23 ANSWER 90 OF 280 CAPLUS COPYRIGHT 1999 ACS

1988:201621 CAPLUS AN

108:201621 DN

Plasmid curing in Staphylococcus aureus by antibiotics affecting the

bacterial cell w

- L23 ANSWER 91 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1989:570481 CAPLUS
- DN 111:170481
- TI Extraction of chromosomal DNA from Staphylococcus and Listeria by a rapid achomopeptidase method
- L23 ANSWER 92 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1988:411930 BIOSIS
- TI LYSOSTAPHIN EFFICACY FOR TREATMENT OF STAPHYLOCOCCUS
  -AUREUS INTRAMAMMARY INFECTION.
- L23 ANSWER 93 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1988:305723 BIOSIS
- TI CORRELATION BETWEEN DNA BASE COMPOSITION AND ROUTINE TESTS FOR THE IDENTIFICATION OF MICROCOCCACEAE ISOLATED FROM SHEEP'S MILK CHEESE.
- L23 ANSWER 94 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 41
- AN 1988:354376 BIOSIS
- TI A DYE RELEASE ASSAY FOR DETERMINATION OF LYSOSTAPHIN ACTIVITY.
- L23 ANSWER 95 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 42
- AN 1988:361059 BIOSIS
- TI THE USE OF A MULTIPOINT INOCULATION METHOD TO PERFORM LYSOSTAPHIN LYSOZYME AND GLYCEROL-ERYTHROMYCIN TESTS FOR THE DIFFERENTIATION OF STAPHYLOCOCCI AND MICROCOCCI.
- L23 ANSWER 96 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 43
- AN 1988:505344 BIOSIS
- TI USE OF LYSOSTAPHIN AND BACITRACIN SUSCEPTIBILITY FOR ROUTINE PRESUMPTIVE IDENTIFICATION OF STAPHYLOCOCCI OF BOVINE ORIGIN.
- L23 ANSWER 97 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1988:449617 CAPLUS
- DN 109:49617
- TI Lysostaphin, construction of its expression plasmids, and its manufacture with Escherichia and Bacillus
- L23 ANSWER 98 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 44
- AN 1987:401901 BIOSIS
- TI ISOLATION OF STOMATOCOCCUS-MUCILAGINOSUS FROM DRUG USER WITH ENDOCARDITIS.
- L23 ANSWER 99 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 45
- AN 1987:209525 BIOSIS
- TI CLONING SEQUENCE AND EXPRESSION OF THE LYSOSTAPHIN GENE FROM STAPHYLOCOCCUS-SIMULANS.
- L23 ANSWER 100 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 46
- AN 1987:229890 BIOSIS
- TI ANALYSIS BY GEL ELECTROPHORESIS WESTERN BLOT AND PEPTIDE MAPPING OF PROTEIN A HETEROGENEITY IN STAPHYLOCOCCUS-AUREUS STRAINS.
- L23 ANSWER 101 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 47
- AN 1988:8625 BIOSIS
- THE MOLECULAR ORGANIZATION OF THE LYSOSTAPHIN GENE AND ITS SEQUENCES REPEATED IN TANDEM.
- L23 ANSWER 102 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS

. . . . .

- AN 1987:240882 BIOSIS
- TI COMPARISON OF 5 METHODS OF DIFFERENTIATING STAPHYLOCOCCUS FROM MICROCOCCUS.



- L23 ANSWER 103 OF 28 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1988:287212 BIO
- TI BACTERICIDAL ACTIVITY OF BLOOD PLATELETS ITS DETERMINATION AND NORMAL VALUES.
- L23 ANSWER 104 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 48
- AN 1987:470749 BIOSIS
- TI PLASMID-ENCODED LYSOSTAPHIN ENDOPEPTIDASE GENE OF STAPHYLOCOCCUS-SIMULANS BIOVAR STAPHYLOLYTICUS.
- L23 ANSWER 105 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 49
- AN 1987:145629 BIOSIS
- TI LYSOSTAPHIN LYSIS PROCEDURE FOR DETECTION OF STAPHYLOCOCCUS-AUREUS BY THE FIREFLY BIOLUMINESCENT ATP METHOD.
- L23 ANSWER 106 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 50
- AN 1986:316342 BIOSIS
- TI RAPID LYSOSTAPHIN TEST TO DIFFERENTIATE STAPHYLOCOCCUS
  AND MICROCOCCUS SPECIES.
- L23 ANSWER 107 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 51
- AN 1987:82789 BIOSIS
- TI LYSOSTAPHIN-BASED ASSAY OF HUMAN GRANULOCYTE FUNCTIONS A REEVALUATION.
- L23 ANSWER 108 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1987:206810 BIOSIS
- TI SYSTEM FOR TESTING THE PHAGOCYTIC CAPACITY OF HUMAN BLOOD PLATELETS.
- L23 ANSWER 109 OF 280 MEDLINE
- AN 86318215 MEDLINE
- [Differentiation between **Staphylococcus** and Micrococcus genera in the routine laboratory diagnosis using the **lysostaphin** sensitivity test].

  Die Differenzierung zwischen den Genera **Staphylococcus** und Micrococcus im Routinelaboratorium mit Hilfe des Tests auf **Lysostaphin**-Empfindlichkeit.
- L23 ANSWER 110 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1986:252980 BIOSIS
- TI A RAPID SIMPLE LYSOSTAPHIN SUSCEPTIBILITY TEST TO DIFFERENTIATE STAPHYLOCOCCI FROM MICROCOCCI.
- L23 ANSWER 111 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 52
- AN 1987:168747 BIOSIS
- TI COMPARISON OF METHODS FOR ROUTINE SEPARATION OF COAGULASE-NEGATIVE STAPHYLOCOCCI FROM MICROCOCCI ISOLATED FROM SHEEP.
- L23 ANSWER 112 OF 280 MEDLINE
- AN 87104076 MEDLINE
- TI Comparison of methods for routine separation of coagulase-negative staphylococci from micrococci isolated from sheep.
- L23 ANSWER 113 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 53
- AN 1986:283577 BIOSIS
- TI INHIBITION OF WALL AUTOLYSIS OF STAPHYLOCOCCI BY SODIUM POLYANETHOLE SULFONATE LIQUOID.
- L23 ANSWER 114 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 54
- AN 1986:458595 BIOSIS
- TI RAPID SEPARATION AND QUANTITATION OF MIXED MICROORGANISMS BY FILTRATION AND BIOLUMINESCENCE.
- L23 ANSWER 115 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS

- AN 1986:372248 BIOSES
  TI ENHANCED PHAGOCY IS OF BACTERIA BY HUMAN NEUTRO ILS FOLLOWING STIMULATION WITH GRANULOCYTE-MACROPHAGE COLONY STIMULATING FACTOR.
- L23 ANSWER 116 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 55
- AN 1987:27757 BIOSIS
- TI EFFECT OF THE COMPOSITION OF REVERSION MEDIUM ON CHANGE OF STAPHYLOCOCCUS-AUREUS LYSOSTAPHIN PROTOPLASTS TO COCCAL FORMS AND L-FORMS.
- L23 ANSWER 117 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 56
- AN 1986:238691 BIOSIS
- TI EVALUATION OF A RAPID TUBE LYSOSTAPHIN TEST TO DIFFERENTIATE BETWEEN STAPHYLOCOCCI AND MICROCOCCI.
- L23 ANSWER 118 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.DUPLICATE 57
- AN 86065463 EMBASE
- TI The contribution of a capsule to survival of staphylococci within bovine neutrophils.
- L23 ANSWER 119 OF 280 MEDLINE
- AN 86199323 MEDLINE
- TI Effect of the composition of the reversion medium on the transformation of coccal forms and L-form of **Staphylococcus** aureus **lysostaphin** protoplasts.
- L23 ANSWER 120 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.DUPLICATE 58
- AN 85077656 EMBASE
- TI Adherence of lysostaphin to and penetration into human monocytes.
- L23 ANSWER 121 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1985:422001 BIOSIS
- TI ANTAGONISTIC ACTIVITIES OF COAGULASE-POSITIVE STAPHYLOCOCCI.
- L23 ANSWER 122 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 59
- AN 1986:214111 BIOSIS
- TI CELL FUSION BETWEEN L-FORMS AND PROTOPLASTS OF STAPHYLOCOCCUS-AUREUS.
- L23 ANSWER 123 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 60
- AN 1985:298045 BIOSIS
- TI DECOMPLEMENTATION ANTIGEN A POSSIBLE DETERMINANT OF STAPHYLOCOCCAL PATHOGENICITY.
- L23 ANSWER 124 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1986:166521 BIOSIS
- TI SPECIES DISTRIBUTION OF COAGULASE-NEGATIVE STAPHYLOCOCCI ISOLATED FROM CLINICAL SOURCES.
- L23 ANSWER 125 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1985:417289 BIOSIS
- TI RAPID CLASSIFICATION OF STAPHYLOCOCCI BY USING LYSOSTAPHIN SENSITIVITY.
- L23 ANSWER 126 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 61
- AN 1984:331781 BIOSIS
- TI INTRA LEUKOCYTIC SEQUESTRATION AS A CAUSE OF PERSISTENT STAPHYLOCOCCUS-AUREUS PERITONITIS IN CONTINUOUS AMBULATORY PERITONEAL DIALYSIS.
- L23 ANSWER 127 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 62
- AN 1984:315916 BIOSIS
- TI COMPARISON OF VARIOUS METHODS FOR DIFFERENTIATION OF STAPHYLOCOCCI AND MICROCOCCI.

AN 1984:172776 CAPLUS

DN 100:172776

- TI Determination of phagocytosis of 32P-labeled Staphylococcus aureus by bovine polymorphonuclear leukocytes
- L23 ANSWER 129 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 63

AN 1984:290332 BIOSIS

- TI STAPHYLOCOCCI ISOLATED FROM ABSCESSES IN SLAUGHTERED ANIMALS CHARACTERIZATION AND EPIDEMIOLOGICAL STUDIES.
- L23 ANSWER 130 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS

AN 1984:291850 BIOSIS

- TI FLUORESCENT STAINING OF INTRA CELLULAR AND EXTRACELLULAR BACTERIA IN BLOOD.
- L23 ANSWER 131 OF 280 SCISEARCH COPYRIGHT 1999 ISI (R)

AN 84:292621 SCISEARCH

- TI LYSOSTAPHIN-DISC-TEST FOR RAPID DIFFERENTIATION OF STAPHYLOCOCCI AND MICROCOCCI
- L23 ANSWER 132 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 64

AN 1984:109874 BIOSIS

- TI IMPROVEMENT IN THE LIPID EXTRACTION OF STAPHYLOCOCCAL CELLS BY LYSOSTAPHIN TREATMENT.
- L23 ANSWER 133 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 65

AN 1985:307468 BIOSIS

- TI A MODIFIED MEDIUM FOR THE RECOVERY OF STAPHYLOCOCCUS FROM WATER.
- L23 ANSWER 134 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS

AN 1984:114373 BIOSIS

- TI A COMPARATIVE STUDY OF METHODS USED FOR THE DIFFERENTIATION AND CHARACTERIZATION OF THE MICROCOCCACEAE.
- L23 ANSWER 135 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS

AN 1985:422018 BIOSIS

- TI COMPARATIVE STUDY OF 4 PROCEDURES FOR SEPARATING STAPHYLOCOCCI FROM MICROCOCCI.
- L23 ANSWER 136 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.DUPLICATE 66

AN 84035605 EMBASE

- TI Concurrent estimation of the kinetics of adhesion and ingestion of Staphylococcus aureus by human polymorphonuclear leukocytes (PMNs).
- L23 ANSWER 137 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 67

AN 1984:197346 BIOSIS

- TI FIBRONECTIN RECEPTORS FROM STAPHYLOCOCCUS-AUREUS.
- L23 ANSWER 138 OF 280 MEDLINE

AN 84067530 MEDLINE

- TI Induction of L-phase variant from protoplast of Staphylococcus aureus.
- L23 ANSWER 139 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 68

AN 1983:282097 BIOSIS

- TI ANTIBIOTIC ACTIVITY AGAINST INTRA LEUKOCYTIC STAPHYLOCOCCUS-AUREUS IN-VITRO AND IN EXPERIMENTAL MASTITIS IN MICE.
- L23 ANSWER 140 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1984:48402 CAPLUS

DN 100:48402

TI Cationic polyelectrolytes activate autolytic wall enzymes in

Staphylococcus are us: modulation by anionic polyelectrolytes in returnation to the survival bacterial constituents in tiss

- L23 ANSWER 141 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 69
- AN 1984:308 BIOSIS
- TI CLASSIFICATION AND LYSOSTAPHIN SUSCEPTIBILITY OF AIRBORNE STAPHYLOCOCCI.
- L23 ANSWER 142 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.
- AN 83212247 EMBASE
- TI [Serum antibody assay to cell wall of Staphylococcus aureus by a hemagglutination test].

  TITRAGE DES ANTICORPS SERIQUES ANTI-PEPTIDOGLYCANE PARIETAL DE STAPHYLOCOCCUS AUREUS PAR HEMAGGLUTINATION INDIRECTE.
- L23 ANSWER 143 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.DUPLICATE 70
- AN 83208111 EMBASE
- TI The influence of lysostaphin on phagocytosis, intracellular bactericidal activity, and chemotaxis of human polymorphonuclear cells.
- L23 ANSWER 144 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1984:317913 BIOSIS
- TI STAPHYLOCOCCAL LIPASE INTRA CELLULAR ENZYME PRODUCTION.
- L23 ANSWER 145 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 71
- AN 1983:246438 BIOSIS
- TI RAPID IDENTIFICATION OF STAPHYLOCOCCUS-AUREUS AND STREPTOCOCCUS-PNEUMONIAE FROM BLOOD CULTURES.
- L23 ANSWER 146 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 72
- AN 1982:275012 BIOSIS
- TI FIBRONECTIN MEDIATES ATTACHMENT OF STAPHYLOCOCCUS-AUREUS TO HUMAN NEUTROPHILS.
- L23 ANSWER 147 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1982:196440 CAPLUS
- DN 96:196440
- TI pH-dependent penicillin tolerance may protect intraleukocytic Staphylococcus aureus from killing by cloxacillin
- L23 ANSWER 148 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 73
- AN 1982:279693 BIOSIS
- TI HIGH LEVEL POTENTIATION OF LYSOSTAPHIN ANTI STAPHYLOCOCCAL ACTIVITY BY LYSOZYME.
- L23 ANSWER 149 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1983:160623 BIOSIS
- TI THE USE OF LYSOSTAPHIN IN IN-VITRO ASSAYS OF PHAGOCYTE FUNCTION ADHERENCE TO AND PENETRATION INTO GRANULOCYTES.
- L23 ANSWER 150 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1983:58533 BIOSIS
- TI IMPROVED METHODOLOGY FOR DETERMINATION OF PHAGOCYTOSIS OF PHOSPHORUS-32 LABELED STAPHYLOCOCCUS-AUREUS BY POLYMORPHONUCLEAR LEUKOCYTES.
- L23 ANSWER 151 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 74
- AN 1983:173412 BIOSIS
- TI IDENTIFICATION OF STAPHYLOCOCCUS-STAPHYLOLYTICUS NRRL-B-2628 AS A BIOVAR OF STAPHYLOCOCCUS-SIMULANS.
- L23 ANSWER 152 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 75
- AN 1983:246419 BIOSIS
- TI BIOLOGIC ACTIVITY OF CELL ASSOCIATED STAPHYLOCOCCAL ENTERO TOXIN A.

- L23 ANSWER 153 OF 2 CAPLUS COPYRIGHT 1999 ACS
- AN 1983:119200 CAP 30
- DN 98:119200
- TI The effect of minocycline and lysostaphin on the intracellular killing of Staphylococcus aureus by polymorphonuclear leukocytes
- L23 ANSWER 154 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 76
- AN 1981:289399 BIOSIS
- TI LYSOSTAPHIN DISC TEST FOR ROUTINE PRESUMPTIVE IDENTIFICATION OF STAPHYLOCOCCI.
- L23 ANSWER 155 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 77
- AN 1982:150958 BIOSIS
- TI SENSITIVITY TO LYSOSTAPHIN AS A CRITERION FOR IDENTIFICATION OF STAPHYLOCOCCI FROM ANIMAL ORIGIN.
- L23 ANSWER 156 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 78
- AN 1982:174713 BIOSIS
- TI PROTEIN A ACTIVITY OF STAPHYLOCOCCUS-HYICUS IN COMPARISON TO PROTEIN A OF STAPHYLOCOCCUS-AUREUS.
- L23 ANSWER 157 OF 280 MEDLINE
- AN 82087003 MEDLINE
- TI [Protein A-activity of Staphylococcus hyicus in comparison to protein A of Staphylococcus aureus (author's transl)].

  Protein A-Aktivitat von Staphylococcus hyicus im Vergleich zu Protein A von Staphylococcus aureus.
- L23 ANSWER 158 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1981:295943 BIOSIS
- TI A POLYMORPHONUCLEAR LEUKOCYTE MONO LAYER SYSTEM FOR STUDIES OF PHAGOCYTOSIS.
- L23 ANSWER 159 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1981:232864 BIOSIS
- TI ENDOGENOUS PYROGEN PRODUCTION BY HUMAN BLOOD MONOCYTES STIMULATED BY STAPHYLOCOCCAL CELL WALL COMPONENTS.
- L23 ANSWER 160 OF 280 MEDLINE
- AN 82172614 MEDLINE
- TI [Sensitivity to lysostaphin lysis of staphylococci isolated from ovine mastitic milk].

  Sensibilidad a la lisis por lisostafina en estafilococos aislados de leche mamitica de oveja.
- L23 ANSWER 161 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1980:67565 BIOSIS
- TI CHARACTERISTICS OF COAGULASE NEGATIVE STAPHYLOCOCCI ISOLATED FROM MARROW TRANSPLANT PATIENTS.
- L23 ANSWER 162 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1980:57229 BIOSIS
- TI EVALUATION OF 6 CORRELATES OF THE COAGULASE TEST FOR IDENTIFYING STAPHYLOCOCCUS-AUREUS.
- L23 ANSWER 163 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1980:211630 CAPLUS
- DN 92:211630
- TI Recombinant plasmids carrying promoters, genes and the origin of DNA replication of the early region of bacteriophage T7
- L23 ANSWER 164 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 79

- AN 1981:190011 BICKIS
  TI MONOCYTES IN IN MATORY BOWEL DISEASE PHAGOCYT S AND INTRA CELLULAR KILLING.
- L23 ANSWER 165 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 80
- AN 1981:140921 BIOSIS
- TI FACILE PENETRATION OF THE STAPHYLOCOCCUS-AUREUS CAPSULE BY LYSOSTAPHIN.
- L23 ANSWER 166 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1980:251516 BIOSIS
- TI ANTIBODY INHIBITION OF POLYMORPHONUCLEAR PHAGOCYTOSIS DISSOCIATION OF BACTERIAL ATTACHMENT AND BACTERIAL KILLING.
- L23 ANSWER 167 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 81
- AN 1981:169147 BIOSIS
- TI RAPID IDENTIFICATION OF STAPHYLOCOCCUS-AUREUS BY USING LYSOSTAPHIN SENSITIVITY.
- L23 ANSWER 168 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 82
- AN 1981:16018 BIOSIS
- TI EFFECT OF VARIOUS BLOOD CULTURE MEDIA ON LYSOSTAPHIN SENSITIVITY OF STAPHYLOCOCCI.
- L23 ANSWER 169 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1980:238746 BIOSIS
- TI CHARACTERIZATION OF STAPHYLOCOCCI ISOLATED FROM MASTITIC COWS IN SPAIN.
- L23 ANSWER 170 OF 280 MEDLINE
- AN 81071179 MEDLINE
- TI The characteristics of extracellular protein secretion by Staphylococcus staphylolyticus.
- L23 ANSWER 171 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1981:47851 BIOSIS
- TI ANTIBODY INHIBITION OF POLYMORPHONUCLEAR PHAGOCYTOSIS DISSOCIATION OF BACTERIAL ATTACHMENT AND BACTERIAL KILLING.
- L23 ANSWER 172 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.
- AN 80197299 EMBASE
- TI Involvement of the cell envelope in plasmid maintenance: plasmid curing during the regeneration of protoplasts.
- L23 ANSWER 173 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 83
- AN 1981:151915 BIOSIS
- TI METHOD TO PROVE INGESTION OF PARTICLES BY MACROPHAGES WITH LIGHT MICROSCOPY.
- L23 ANSWER 174 OF 280 MEDLINE
- AN 81126174 MEDLINE
- TI Method to prove investigation of particles by macrophages with light microscopy.
- L23 ANSWER 175 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 84
- AN 1981:294063 BIOSIS
- TI SELECTION IN-VITRO OF ANTIBIOTICS WITH ACTIVITY AGAINST INTRA CELLULAR STAPHYLOCOCCUS-AUREUS.
- L23 ANSWER 176 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 85
- AN 1981:131177 BIOSIS
- TI THE EFFECTS OF CLOXACILLIN ON STAPHYLOCOCCI STAPHYLOCOCCUS-AUREUS PHAGOCYTOSED BY BOVINE NEUTROPHILS.

- L23 ANSWER 177 OF 28 BIOSIS COPYRIGHT 1999 BIOSIS
- 1980:5431 BIOSI AN
- C STUDY OF CELL WALL PROTEIN OF STAPHYLOCOCCUS-AUREUS A KIN ΤI RELEASE BY LYSOSTAPHIN.
- L23 ANSWER 178 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 86
- 1979:228510 BIOSIS AN
- RELATIONSHIP BETWEEN LYSOSTAPHIN ENDO PEPTIDASE PRODUCTION AND TI CELL WALL COMPOSITION IN STAPHYLOCOCCUS-STAPHYLOLYTICUS.
- L23 ANSWER 179 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 87
- 1980:119320 BIOSIS AN
- RAPID SCREENING TEST FOR STAPHYLOCOCCUS-AUREUS USING TΙ LYSOSTAPHIN.
- L23 ANSWER 180 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- 1980:47072 BIOSIS ΑN
- PHAGOCYTIC POTENTIAL OF HAIRY CELLS. ΤI
- L23 ANSWER 181 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- 1980:48569 BIOSIS ΑN
- NORMAL NEUTROPHIL PHAGOCYTIC AND BACTERICIDAL ACTIVITY IN EXPERIMENTAL NUTRITIONAL IRON DEFICIENCY.
- L23 ANSWER 182 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 88
- 1980:149352 BIOSIS AN
- PHAGOCYTIC POTENTIAL OF HAIRY CELLS. ΤI
- L23 ANSWER 183 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 89
- 1979:252617 BIOSIS ΑN
- HAIRY CELL LEUKEMIA A BONE MARROW DERIVED CELL LYMPHOCYTIC DISORDER. TТ
- L23 ANSWER 184 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.DUPLICATE 90
- 79175285 EMBASE AN
- Evaluation of the lysostaphin-susceptibility test for the TТ classification of staphylococci.
- L23 ANSWER 185 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- 1979:89585 BIOSIS NΑ
- AGE RELATED CHANGES IN PULMONARY MACROPHAGES. ΤI
- L23 ANSWER 186 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 91
- 1979:155938 BIOSIS AN
- COUNTER IMMUNO ELECTROPHORETIC DETECTION OF A HIGH INCIDENCE OF PRECIPITIN REACTIONS IN NORMAL HUMAN SERA AGAINST STAPHYLOCOCCAL TEICHOIC ACIDS AND PROTEIN A.
- L23 ANSWER 187 OF 280 MEDLINE

DUPLICATE 92

- 79164149 MEDLINE
- Cell-wall proteins of Staphylococcus aureus : a kinetic study of ΤI release by lysostaphin.
- L23 ANSWER 188 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 93
- 1979:155910 BIOSIS AΝ
- USE OF LYSOSTAPHIN TO REMOVE CELL ADHERENT STAPHYLOCOCCI TТ DURING IN-VITRO ASSAYS OF PHAGOCYTE FUNCTION.
- L23 ANSWER 189 OF 280 SCISEARCH COPYRIGHT 1999 ISI (R)
- 78:236709 SCISEARCH AΝ
- USE OF LYSOSTAPHIN TO REMOVE CELL-ADHERENT STAPHYLOCOCCI ΤI DURING INVITRO ASSAYS OF PHAGOCYTE FUNCTION
- L23 ANSWER 190 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS



- AN 1979:252140 BIOSIS
  TI EVALUATION OF THE LYSOSTAPHIN SUSCEPTIBILITY TESTS OR THE CLASSIFICATION STAPHYLOCOCCI.
- L23 ANSWER 191 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1978:100654 BIOSIS
- TI USE OF LYSOSTAPHIN TO REMOVE CELL ADHERENT STAPHYLOCOCCI DURING IN-VITRO ASSAYS OF PHAGOCYTE FUNCTION.
- L23 ANSWER 192 OF 280 MEDLINE
- AN 78041466 MEDLINE
- TI Comparison of fatty acid composition of stable L-phase variants of Staphylococcus aureus induced by three differnet mechanisms.
- L23 ANSWER 193 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1977:418924 CAPLUS
- DN 87:18924
- TI Gentle lysis of Staphylococcus aureus at low temperature
- L23 ANSWER 194 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 94
- AN 1977:224543 BIOSIS
- TI RAPID SOLID PHASE RADIOASSAY FOR STAPHYLOCOCCAL PROTEIN A.
- L23 ANSWER 195 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1977:498718 CAPLUS
- DN 87:98718
- TI Coagulase-negative staphylococci
- L23 ANSWER 196 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1977:174210 BIOSIS
- TI KINETICS OF STAPHYLOCOCCAL OPSONIZATION ATTACHMENT INGESTION AND KILLING BY HUMAN POLYMORPHONUCLEAR LEUKOCYTES A QUANTITATIVE ASSAY USING TRITIATED THYMIDINE LABELED BACTERIA.
- L23 ANSWER 197 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1978:142383 BIOSIS
- TI LYSOSTAPHIN ENDO PEPTIDASE CATALYZED TRANS PEPTIDATION REACTIONS OF THE IMINO TRANSFER TYPE.
- L23 ANSWER 198 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 95
- AN 1977:223414 BIOSIS
- TI EVALUATION OF PHAGOCYTOSIS OF STAPHYLOCOCCUS-AUREUS WITH THE AID OF LYSOSTAPHIN.
- L23 ANSWER 199 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1978:71625 BIOSIS
- TI CELL WALL COMPOSITION AND LYSOSTAPHIN ENDO PEPTIDASE RESISTANCE IN STAPHYLOCOCCUS-STAPHYLOLYTICUS.
- L23 ANSWER 200 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.DUPLICATE 96
- AN 77166192 EMBASE
- TI The occurrence of cell associated enterotoxin B in Staphylococcus aureus.
- L23 ANSWER 201 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1977:66529 CAPLUS
- DN 86:66529
- TI Inhibition of leukocyte migration by peptidoglycan fragments
- L23 ANSWER 202 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.
- AN 77097184 EMBASE
- TI The rate of phagocytosis and killing of staphylococci in the murine lung.
- L23 ANSWER 203 OF 280 MEDLINE

- AN 77114939 MEDLINE
  TI The identification of staphylococci in clinical food microbiology laboratories.
- L23 ANSWER 204 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.
- AN 77192012 EMBASE
- TI Susceptibility of **staphylococci** of various cell wall structures to **lysostaphin** and its separated enzymes.
- L23 ANSWER 205 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1977:84175 CAPLUS
- DN 86:84175
- TI Susceptibility of staphylococci of various cell wall structure to lysostaphin and its separated enzymes
- L23 ANSWER 206 OF 280 SCISEARCH COPYRIGHT 1999 ISI (R)
- AN 77:62835 SCISEARCH
- TI SUSCEPTIBILITY OF STAPHYLOCOCCI OF VARIOUS CELL-WALL STRUCTURE TO LYSOSTAPHIN AND ITS SEPARATED ENZYMES
- L23 ANSWER 207 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.
- AN 77097391 EMBASE
- TI Effect of in vitro exposure to ethanol on the antibacterial activity of alveolar macrophages in pulmonary lavage fluid.
- L23 ANSWER 208 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 97
- AN 1976:185044 BIOSIS
- TI MEASUREMENT OF PHAGOCYTOSIS OF PHOSPHORUS-32 LABELED STAPHYLOCOCCUS-AUREUS BY BOVINE LEUKOCYTES LYSOSTAPHIN DIGESTION AND INHIBITORY EFFECT OF CREAM.
- L23 ANSWER 209 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 98
- AN 1976:154005 BIOSIS
- TI INHIBITION BY GLUTARALDEHYDE OF LYSOSTAPHIN INDUCED LYSIS OF STAPHYLOCOCCUS-AUREUS.
- L23 ANSWER 210 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1975:147241 BIOSIS
- TI DENSITY GRADIENT SEPARATION OF LYMPHOID CELLS ADHERING TO PROTEIN A CONTAINING STAPHYLOCOCCI.
- L23 ANSWER 211 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.
- AN 75047208 EMBASE
- TI The role of peroxidase in the bactericidal activity of human blood eosinophils.
- L23 ANSWER 212 OF 280 EMBASE COPYRIGHT 1999 ELSEVIER SCI. B.V.DUPLICATE 99
- AN 75037313 EMBASE
- TI Lack of correlation between methicillin resistance and susceptibility to lysostaphin in Staphylococcus aureus.
- L23 ANSWER 213 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 100
- AN 1975:1172 BIOSIS
- TI SYSTEMIC LYSOSTAPHIN IN MAN APPARENT ANTI MICROBIAL ACTIVITY IN A NEUTROPENIC PATIENT.
- L23 ANSWER 214 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1974:566102 CAPLUS
- DN 81:166102
- TI Amino acid transport and staphylococcal membrane vesicles
- L23 ANSWER 215 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1974:159590 BIOSIS

- TI INTERACTION OF I RA LEUKOCYTIC BACTERIA AND ANTIBIO.....
- L23 ANSWER 216 OF 28 CAPLUS COPYRIGHT 1999 ACS
- AN 1974:11679 CAPLUS
- DN 80:11679
- TI Lysostaphin. Separation and characterization of three enzymes
- L23 ANSWER 217 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 101
- AN 1974:164644 BIOSIS
- TI SUSCEPTIBILITY OF STAPHYLOCOCCUS-EPIDERMIDIS TO LYSOSTAPHIN AND MAJOR ANTI MICROBIAL AGENTS.
- L23 ANSWER 218 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1974:402168 CAPLUS
- DN 81:2168
- TI Structure and immunology of protein A
- L23 ANSWER 219 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1973:234741 BIOSIS
- TI CULTURAL PHAGOCYTIC AND BACTERICIDAL CHARACTERISTICS OF PERITONEAL MACROPHAGES.
- L23 ANSWER 220 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1973:98797 BIOSIS
- TI L FORMS.
- L23 ANSWER 221 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1974:518533 CAPLUS
- DN 81:118533
- TI Lysostaphin by fermentation
- L23 ANSWER 222 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 102
  - AN 1973:139621 BIOSIS
- TI PROTEIN A ISOLATED FROM STAPHYLOCOCCUS-AUREUS AFTER DIGESTION WITH LYSOSTAPHIN.
- L23 ANSWER 223 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 103
- AN 1973:161704 BIOSIS
- TI BINDING SITES FOR CATIONIC PROTEINS ON STAPHYLOCOCCI.
- L23 ANSWER 224 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1973:414468 CAPLUS
- DN 79:14468
- TI Lysostaphin. Model for a specific enzymic approach to infectious disease
- L23 ANSWER 225 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1973:156024 BIOSIS
- TI LOCALIZATION OF PROTEIN A IN THE BACTERIA.
- L23 ANSWER 226 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1971:497260 CAPLUS
- DN 75:97260
- TI Lysostaphin fermentation with accelerated time cycle
- L23 ANSWER 227 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 104
- AN 1971:204344 BIOSIS
- TI NUCLEASE PRODUCTION AND LYSOSTAPHIN SUSCEPTIBILITY OF STAPHYLOCOCCUS-AUREUS AND OTHER CATALASE POSITIVE COCCI.
- L23 ANSWER 228 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 105
- AN 1972:118261 BIOSIS
- TI EFFICACY AND SAFETY OF TOPICAL LYSOSTAPHIN TREATMENT OF PERSISTENT NASAL CARRIAGE OF STAPHYLOCOCCUS-AUREUS.

 $\sqrt{}$ 

- L23 ANSWER 229 OF 2 CAPLUS COPYRIGHT 1999 ACS
- AN 1971:95938 CAR
- DN 74:95938
- TI Cycloserine induction, propagation, and antimicrobial susceptibility of wall-defective Staphylococcus aureus
- L23 ANSWER 230 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1971:515798 CAPLUS
- DN 75:115798
- Modified assay of neutrophil function. Use of lysostaphin to differentiate defective phagocytosis from impaired intracellular killing
- L23 ANSWER 231 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 106
- AN 1971:199731 BIOSIS
- TI STUDIES ON ENDO BETA-N ACETYL GLUCOSAMINIDASE STAPHYLOLYTIC PEPTIDASE AND N ACETYLMURAMYL L ALANINE AMIDASE IN LYSOSTAPHIN AND FROM STAPHYLOCOCCUS-AUREUS.
- L23 ANSWER 232 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1971:495754 CAPLUS
- DN 75:95754
- TI Autolytic activity in methicillin-resistant Staphylococcus aureus
- L23 ANSWER 233 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1971:83621 BIOSIS
- TI BACTERIO PHAGE REPRODUCTION IN LYSOSTAPHIN TREATED STAPHYLOCOCCUS-AUREUS 44-A-HJD.
- L23 ANSWER 234 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1972:191909 BIOSIS
- TI ENDOGENOUS RESERVE OF STAPHYLOCOCCI.
- L23 ANSWER 235 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1972:174588 BIOSIS
- TI LYSIS AND LYSATES STUDY WITH STAPHYLOCOCCI.
- L23 ANSWER 236 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 107
- AN 1971:207550 BIOSIS
- TI LIGHT MICROSCOPY AND SCANNING BEAM ELECTRON MICROSCOPY OF WALL DEFECTIVE STAPHYLOCOCCUS-AUREUS INDUCED BY LYSOSTAPHIN.
- L23 ANSWER 237 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 108
- AN 1971:111924 BIOSIS
- TI MOLECULAR PROPERTIES OF LYSOSTAPHIN A BACTERIOLYTIC AGENT SPECIFIC FOR STAPHYLOCOCCUS-AUREUS.
- L23 ANSWER 238 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1971:30379 BIOSIS
- TI LIGHT MICROSCOPE AND SCANNING BEAM ELECTRON MICROSCOPY OF WALL DEFECTIVE STAPHYLOCOCCUS-AUREUS INDUCED BY LYSOSTAPHIN.
- L23 ANSWER 239 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1970:52190 CAPLUS
- DN 72:52190
- TI Effects of lysostaphin and its two active components on stable wall-defective forms of Staphylococcus aureus
- L23 ANSWER 240 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1970:516447 CAPLUS
- DN 73:116447
- TI Characterization of a Staphylococcus aureus bacteriocin

L23 ANSWER 241 OF 28 BIOSIS COPYRIGHT 1999 BIOSIS DUFLICATE 103

AN 1970:198517 BI

TI LYSOSTAPHIN IND ED OSMOTICALLY FRAGILE STAPHYLO CCUS
-AUREUS CELLS.

- L23 ANSWER 242 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1971:431837 CAPLUS

DN 75:31837

- TI Cell walls of methicillin-resistant Staphylococcus aureus
- L23 ANSWER 243 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 110

AN 1969:121678 BIOSIS

- TI INST ELECTRON MICROSCOPY AND VIABILITY OF ENZ LYSOSTAPHIN ANTI INFECT INDUCED STAPHYLOCOCCAL SPHEROPLASTS PROTOPLAST-LIKE BODIES AND PROTOPLASTS STAPHYLOCOCCUS-AUREUS.
- L23 ANSWER 244 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1970:97622 CAPLUS

DN 72:97622

- TI Staphylococcal spheroplasts and L colonies. IV. Antimicrobial susceptibility of stable methicillin-induced and lysostaphin -induced spheroplasts
- L23 ANSWER 245 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1969:59021 BIOSIS
- TI TREATMENT OF CANINE STAPHYLOCOCCAL ENDO CARDITIS WITH ENZ LYSOSTAPHIN ANTI INFECT OR OXACILLIN ANTI INFECT.
- L23 ANSWER 246 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 111

AN 1971:2741 BIOSIS

- TI THE USE OF LYSOSTAPHIN IN TREATMENT OF STAPHYLOCOCCAL WOUND INFECTIONS.
- L23 ANSWER 247 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 112

AN 1969:155552 BIOSIS

- TI STAPHYLOCOCCAL SPHEROPLASTS AND L COLONIES III INDUCTION BY ENZ LYSOSTAPHIN STAPHYLOCOCCUS-AUREUS.
- L23 ANSWER 248 OF 280 MEDLINE DUPLICATE 113

AN 69137459 MEDLINE

- TI Staphylococcus aureus response to lysostaphin in some fermented foods.
- L23 ANSWER 249 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS

AN 1969:139395 BIOSIS

- TI STAPHYLOCOCCUS-AUREUS RESPONSE TO ENZ LYSOSTAPHIN DISINFECT IN SOME FERMENTED FOODS CHEESE SAUSAGE.
- L23 ANSWER 250 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 114

AN 1969:132965 BIOSIS

- TI TRANSFECTION OF ENZ LYSOSTAPHIN ANTI INFECT TREATED CELLS OF STAPHYLOCOCCUS-AUREUS STAPHYLOCOCCAL PHAGES 53 44A.
- L23 ANSWER 251 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1968:495105 CAPLUS
- DN 69:95105
- TI Lysostaphin production by fermentation
- L23 ANSWER 252 OF 280 MEDLINE

DUPLICATE 115

AN 68406273 MEDLINE

TI Comparative inhibition of methicillin-resistant strains of **Staphylococcus** aureus by **lysostaphin** and other antibiotics.

- L23 ANSWER 253 OF MEDLINE
- AN 68406272
- TI Susceptibility of coagulase-negative staphylococci to lysostaphin and other antibiotics.
- L23 ANSWER 254 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1968:75873 CAPLUS
- DN 68:75873
- TI Use of lysostaphin in the isolation of highly polymerized, deoxyribonucleic acid and in the taxonomy of aerobic Micrococcaceae
- L23 ANSWER 255 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 117
- AN 1969:218124 BIOSIS
- TI OSMOTIC FRAGILITY AND VIABILITY OF ENZ LYSOSTAPHIN ANTI INFECT INDUCED STAPHYLOCOCCAL SPHEROPLASTS STAPHYLOCOCCUS
  -AUREUS
- L23 ANSWER 256 OF 280 MEDLINE
- AN 70001548 MEDLINE
- TI Staphylococcal spheroplasts and L colonies. IV. **Antimicrobial** susceptibility of stable methicillin-induced and **lysostaphin** -induced spheroplasts.
- L23 ANSWER 257 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1969:116858 BIOSIS
- TI IN-VITRO ACTIVITY OF ENZ LYSOSTAPHIN ANTI INFECT BIOASSAY IN SERUM USING STAPHYLOCOCCUS-AUREUS DOG.
- L23 ANSWER 258 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 118
- AN 1969:99990 BIOSIS
- TI ENZ LYSOSTAPHIN AN ENZYMATIC APPROACH TO STAPHYLOCOCCAL DISEASE III COMBINED ENZ LYSOSTAPHIN ANTI INFECT METHICILLIN ANTI INFECT THERAPY OF ESTABLISHED STAPHYLOCOCCAL ABSCESSES IN MICE RENAL.
- L23 ANSWER 259 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1968:425281 CAPLUS
- DN 69:25281
- TI Lysostaphin. II. Sensitivity of 230 Staphylococcus aureus strains of animal origin to lysostaphin
- L23 ANSWER 260 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1968:425280 CAPLUS
- DN 69:25280
- TI Lysostaphin. I. Sensitivity of 355 Staphylococcus aureus strains of human origin to lysostaphin
- L23 ANSWER 261 OF 280 MEDLINE

DUPLICATE 119

- AN 67217944 MEDLINE
- TI Lytic action of lysostaphin on susceptible and resistant strains of Staphylococcus aureus.
- L23 ANSWER 262 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1967:63805 CAPLUS
- DN 66:63805
- TI Immunologically active cell wall peptide polymer of Staphylococcus aureus
- L23 ANSWER 263 OF 280 CAPLUS COPYRIGHT 1999 ACS
- AN 1967:45319 CAPLUS
- DN 66:45319
- TI Lysostaphin in experimental renal infections

1/

\/

L23 ANSWER 264 OF 286 MEDLINE

AN 67205710 ME NE

TI Lysostaphin: an enzymatic approach to staphyloca al disease. II. In vivo studies.

DUPLICATE 121

L23 ANSWER 265 OF 280 MEDLINE

AN 67263769 MEDLINE

TI Lysostaphin: an enzymatic approach to staphylococcal disease. I. In vitro studies.

L23 ANSWER 266 OF 280 MEDLINE

AN 68368241 MEDLINE

TI Effect of lysostaphin on staphylococcal carriage in infants and children.

L23 ANSWER 267 OF 280 CAPLUS COPYRIGHT 1999 ACS

AN 1967:36424 CAPLUS

DN 66:36424

TI Therapeutic activity of lysostaphin in experimental staphylococcal infections

L23 ANSWER 268 OF 280 MEDLINE

AN 68368299 MEDLINE

TI Studies in experimental staphylococcal endocarditis in dogs. VI. Treatment with lysostaphin.

L23 ANSWER 269 OF 280 CAPLUS COPYRIGHT 1999 ACS

AN 1967:452667 CAPLUS

DN 67:52667

TI Selective activity of lysostaphin in vivo

L23 ANSWER 270 OF 280 MEDLINE

AN 67081173 MEDLINE

TI Staphylococcal strains with relation to lysostaphin sensistivity.

L23 ANSWER 271 OF 280 CAPLUS COPYRIGHT 1999 ACS

AN 1967:479901 CAPLUS

DN 67:79901

TI In vitro activity of lysostaphin

L23 ANSWER 272 OF 280 MEDLINE

AN 67055171 MEDLINE

TI Growth inhibition of unusual strains of **Staphylococcus** aureus by **lysostaphin** and other antistaphylococcal antibiotics.

L23 ANSWER 273 OF 280 MEDLINE

AN 67042251 MEDLINE

TI In vitro effect of lysostaphin, neomycin, and bacitracin on Staphylococcus aureus.

L23 ANSWER 274 OF 280 CAPLUS COPYRIGHT 1999 ACS

AN 1967:420242 CAPLUS

DN 67:20242

TI Experimental observations on staphylococcal disease

L23 ANSWER 275 OF 280 MEDLINE

AN 66113613 MEDLINE

TI The role of NaCl in the lysis of **Staphylococcus** aureus by lysostaphin.

L23 ANSWER 276 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS

AN 1996:218739 BIOSIS

- TI Safety assessment of genetically modified microorganisms applied in medefermentations.
- L23 ANSWER 277 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1978:78502 BIOSIS
- TI EFFECT OF LYSOSTAPHIN ENDO PEPTIDASE PRODUCTION ON THE CELL WALL OF STAPHYLOCOCCUS-STAPHYLOLYTICUS.
- L23 ANSWER 278 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1978:77936 BIOSIS
- TI COMPARATIVE EFFECTS OF POLYENE ANTIBIOTICS ON RABBIT ALVEOLAR MACROPHAGES IN-VITRO.
- L23 ANSWER 279 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1969:27404 BIOSIS
- TI STUDIES IN EXPERIMENTAL STAPHYLOCOCCAL ENDO CARDITIS IN DOGS VI TREATMENT WITH ENZ LYSOSTAPHIN ANTI INFECT.
- L23 ANSWER 280 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
- AN 1969:27414 BIOSIS
- TI EFFECT OF ENZ LYSOSTAPHIN ANTI INFECT ON STAPHYLOCOCCAL CARRIAGE IN INFANTS AND CHILDREN STAPHYLOCOCCUS-AUREUS ANTIBODY FORMATION PASSIVE CUTANEOUS SENSITIVITY.

Page 1

02/08/99

M.BORIN

```
(FILE 'USPATFULL, WPIDS, BIOSIS, EMBASE, MEDLINE, CAPLUS, SCISEARCH,
     INVESTEXT, DRUGU' ENTERED AT 11:11:15 ON 08 FEB 1999)
L13
             17 FILE USPATFULL
             12 FILE WPIDS
L14
            162 FILE BIOSIS
L15
            83 FILE EMBASE
L16
            117 FILE MEDLINE
L17
            137 FILE CAPLUS
L18
             65 FILE SCISEARCH
L19
              O FILE INVESTEXT
L20
              6 FILE DRUGU
L21
     TOTAL FOR ALL FILES
            599 S LYSOSTAPHIN (10A) (ANTIMICROB#### OR MICROB### OR STAPHYLOCOC
L22
            280 DUPLICATE REMOVE L15-L19 (284 DUPLICATES REMOVED)
L23
=> d 113 pi,bib, kwic 1-17
L13 ANSWER 1 OF 17 USPATFULL
       US 5858962 990112
       1999:4620 USPATFULL
       Composition for treating mastitis and other staphylococcal infections
       Blackburn, Peter, New York, NY, United States
       Polak, June, Brooklyn, NY, United States
       Ambi Inc., Tarrytown, NY, United States (U.S. corporation)
PA
                                                                         4
       US 5858962 990112
       US 93-168687 931216 (8)
ΑI
       Continuation of Ser. No. US 89-440092, filed on 22 Nov 1989, now
RLI
       abandoned which is a continuation of Ser. No. US 88-188183, filed on 28
       Apr 1988, now abandoned which is a continuation-in-part of Ser. No. US
       87-48412, filed on 11 May 1987, now abandoned
EXNAM Primary Examiner: Weddington, Kevin E.
LREP
       White & Case L.L.P.
       Number of Claims: 14
CLMN
       Exemplary Claim: 1
ECL
       1 Drawing Figure(s); 1 Drawing Page(s)
LN.CNT 733
       Lysostaphin is used to eliminate and cure
AΒ
     staphylococcal infections including the cure of mastitis by
       intramammary infusion. Administration of from 2 mg to 400 mg of
     lysostaphin to an infected bovine mammary gland eliminates
     staphylococci, and the reoccurrence common with antibiotic
       therapy is not observed. Teat-dips containing lysostaphin, mutanolysin
       and lysozyme can be used as.
       This application relates to the use of lysostaphin in the
SUMM
       treatment and prevention of staphylococcal infection and, in
       particular, to the treatment and prevention of staphylococcal bovine
```

```
mastitis.
       Lysostaphin is pacteriocin secreted by a single known strain
SUMM
       of Staphylococcus simulans originally isolated and named
     Staphylococcus staphylolyticus by Schindler and Schuhardt. The
       production of lysostaphin by S. staphylolyticus has been
       described previously in U.S. Pat. No. 3,278,378 issued Oct. 11, 1966 and
       in Proceedings of.
       Bacteriocins are proteins secreted by bacteria that kill and sometimes
SUMM
       lyse related bacteria. For example, lysostaphin lyses and
       kills practically all known staphylococcal species but is
       inactive against bacteria of all other genera. Lysostaphin,
       isolated from culture filtrates of S. simulans (NRRL B-2628) grown
       according to published references, is an endopeptidase which cleaves the
       polyglycine cross-links of the peptidoglycan found in the cell walls of
     staphylococci. In addition, cultures that produce
     lysostaphin appear to be resistant to its activities while
       cultures grown under non-lysostaphin producing conditions are sensitive.
       Studies on the possible mechanism of antibiotic evasion of phagocytized
SUMM
     staphylococci in mastitis treatment show that
     lysostaphin had been rejected as a candidate for destroying
       phagocytized staphylococci. Craven et al., 29 Research in
       Veterinary Science 57 (1980); Craven et al., 21 Antimicrobial Agents and
       Chemotherapy 618 (1982); . . . Comp. Immun. Microbial. Infect. Dis. 447 (1982)) Craven et al., 51 Journal of Dairy Research 513 (1984). In
       these experiments lysostaphin was used in vitro as a
       pretreatment to destroy extracellular staphylococci prior to
       exposing the phagocytized staphylococci to cloxacillin,
       gentamicin or lysostaphin. Craven et al.'s results strongly
       suggest that lysostaphin would have no effect on mastitis
       since intracellular staphylococci were still viable after 20
       hours of incubation in a lysostaphin containing solution. 51
       Journal of Dairy Research at 515-516, and Table 2.
                of chronic nasal staphylococcal infections (Quickel, Jr. et
SUMM
       al., 22 Applied Microbiology 446 (1971)). In one case of a resistant
     staphylococcal infection, lysostaphin was given
       systemically (Stark et al., 291 Medical Intelligence 239 (1974)). In
       general, however, there has been great skepticism and.
             . amount of lysostaphin, with or without surfactant, EDTA,
SUMM
       penicillin or other potentiating agents, are used to achieve elimination
       of the staphylococcal infection. Preferably such infusions
       contain between 2 to 400 mg lysostaphin when no potentiating
       agents are present. In combinations containing potentiating agents, the
       required effective doses of lysostaphin can be lowered.
              . by conventional antibiotic (e.g. penicillin) therapy. In
SUMM
       addition, penicillin and other similar acting substances may also be
       useful together with lysostaphin as an agent against
     staphylococcal infection and contamination.
          . . strain 00 containing a recombinant plasmid which directs the
DETD
       synthesis of lysostaphin, as this provides for both high levels of
     lysostaphin production substantially free from
      staphylococcal immunogenic contaminants and facile
     lysostaphin purification since the lysostaphin
       accumulates directly in the growth medium. Bacillus sphaericus
       transformants containing the plasmid pBC16-1L have been found to be
       particularly suited.
                 developed either chronic or acute staphylococcal bovine
 DETD
       mastitis despite prophylactic treatment. A single dose of from 2 to 400
       mg lysostaphin per milk gland will eliminate the infection and
        cure staphylococcal mastitis in most instances. Additional
        doses of lysostaphin may be indicated where the infection is
       persistent. Doses significantly higher than 400 mg are not recommended
        as they can.
        Table IC demonstrates the synergistic effect of lysostaphin
 DETD
```

09/120030

/penicillin combinations on three strains of statey\_ococc.

Depending on the doses of each, the combination of lysostaphin plus penicillin can be 100 to 1000 times more effective than either lysostaphin or penicillin alone with all three strains.

TABLE ID

DETD

A Comparison of the Effect of the Combination of Lysostaphin and Penicillin Versus Their Sequential Effects on the Survival of Staphylococcus aureus (Strain RN451) in milk at 37.degree. C.

Pen(2h)/ lspn(2h)/

combo(2h) lspr(2h)

pen(2h)

1spn(0.5h)

pen(0.5h)

% survival 0.0005 23 25 0.3 10

DETD

. . . lysostaphin which were sufficient to eliminate the infection did not produce adverse side effects and indicated that intramammary infusions of lysostaphin are effective against

staphylococcal mastitis. At 125 .mu.g/kg, glands were cleared of
 infection by the 6 hour post-treatment sample and remained clear
 throughout the. . .

DETD

TABLE IV

Efficacy of Intramammary Infusion of Lysostaphin Toward Experimental STAPHYLOCOCCAL Mastitis in Guinea Pig

**Lysostaphin** Dose .mu.g/kg
ZERO 1.0 5.0 25.0 62.5 125.0

Number of animals

(0/10) (1/0) (1/2)

(2/2) (1/1)

(7/7)

cleared of
infection

DETD It can be seen from these examples that lysostaphin is effective for treatment of staphylococcal mastitis and that its effect is greatly enhanced when used in combination with penicillin or with substances such as mild. . .

CLM What is claimed is:

1. A composition for killing **staphylococci** comprising **lysostaphin** and an agent which synergistically enhances the bactericidal activity of the lysostaphin, and which is in an amount effective to produce the synergistic enhancement, selected from the group consisting of penicillin, bacitracin, methicillin, cephalosporin and polymyxin and wherein the **lysostaphin** and the agent are together in amounts effective to kill **staphylococci**.

2. A composition for killing staphylococci comprising lysostaphin and at least one agent which synergistically enhances the bactericidal activity of the lysostaphin, and which is in an amount. . .

L13 ANSWER 2 OF 17 USPATFULL PI US 5776712 980707

```
1998:78961 USEATFULL
Methods and materials for the detection Kuusela, Pentti, Helsinki, Finland
ΑN
                                                          ococcus aureus
                        ials for the detection of Stap
ΤI
IN
       Hilden, Pekka, Helsinki, Finland
       Helsinki University Licensing, Ltd., Helsinki, Finland (non-U.S.
PA
       corporation)
                   980707
PΙ
       US 5776712
       US 96-610389 960304 (8)
ΑI
       Continuation-in-part of Ser. No. US 93-169524, filed on 17 Dec 1993, now
RLI
       patented, Pat. No. US 5496706
DT
       Utility
      Primary Examiner: Housel, James C.; Assistant Examiner: Shaver, Jennifer
EXNAM
       Marshall, O'Toole, Gerstein, Murray & Borun
LREP
       Number of Claims: 10
CLMN
       Exemplary Claim: 1
ECL
       4 Drawing Figure(s); 3 Drawing Page(s)
DRWN
LN.CNT 910
       The present invention provides an approximately 230 kDa protein,
SUMM
       designated MRSA-230, which is isolated from lysostaphin
       digests of methicillin-resistant Staphylococcus aureus which
       test negative in standard S. aureus agglutination assays. Thus, the
       invention includes a purified and isolated MRSA-230 protein,.
       A preferred biological sample from which to purify MRSA-230 protein or
SUMM
       fragments is derived from a lysostaphin digest of
     Staphylococcus aureus. Lysostaphin is an enzyme known
       to specifically cleave the pentaglycine bridge which cross-links S.
       aureus surface proteins to the S. aureus.
       Lysostaphin digests of non-agglutinating, methicillin-
DETD
       resistant S. aureus were conducted. Staphylococci grown in 1
       liter Todd-Hewitt medium were suspended in 20 ml NaCl/P.sub.i (0.5M
       sodium phosphate, pH 7.4, 0.14M sodium chloride).
L13 ANSWER 3 OF 17 USPATFULL
       US 5760026 980602
PΤ
       1998:61641 USPATFULL
ΝA
       Method for treating mastitis and other staphylococcal infections
ΤI
       Blackburn, Peter, New York, NY, United States
IN
       Polak, June, Brooklyn, NY, United States
       Ambi Inc., Tarrytown, NY, United States (U.S. corporation)
PA
       US 5760026 980602
PΙ
       US 94-303551 940909 (8)
ΑI
       Continuation of Ser. No. US 92-935121, filed on 20 Aug 1992, now
RLI
       abandoned which is a continuation of Ser. No. US 90-535286, filed on 8
       Jun 1990, now abandoned which is a continuation of Ser. No. US
       88-188183, filed on 28 Apr 1988, now abandoned which is a
       continuation-in-part of Ser. No. US 87-48412, filed on 11 May 1987, now
       abandoned
       Utility
EXNAM Primary Examiner: Weddington, Kevin E.
       White & Case L.L.P.
LREP
       Number of Claims: 5
CLMN
       Exemplary Claim: 1
ECL
        1 Drawing Figure(s); 1 Drawing Page(s)
LN.CNT 844
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
        Lysostaphin is used to eliminate and cure
     staphylococcal infections including the cure of mastitis by
        intramammary infusion. Administration of from 2 mg to 400 mg of
     lysostaphin to an infected bovine mammary gland eliminates
     staphylococci, and the reoccurrence common with antibiotic
        therapy is not observed. Teat-dips containing lysostaphin, mutanolysin
        and lysozyme can be used as.
        This application relates to the use of lysostaphin in the
```

SUMM

```
treatment and prevention of staphylococcal infersion and, in particular, to be treatment and prevention of bhylococcal
                        e treatment and prevention of
                                                          phylococcal bovine
       particular, to
       mastitis.
       Lysostaphin is a bacteriocin secreted by a single known strain
SUMM
       of Staphylococcus simulans originally isolated and named
     Staphylococcus staphylolyticus by Schindler and Schuhardt. The
       production of lysostaphin by S. staphylolyticus has been
       described previously in U.S. Pat. No. 3,278,378 issued Oct. 11, 1966 and
       in Proceedings of.
       Bacteriocins are proteins secreted by bacteria that kill and sometimes
SUMM
       lyse related bacteria. For example, lysostaphin lyses and
       kills practically all known staphylococcal species but is
       inactive against bacteria of all other genera. Lysostaphin,
       isolated from culture filtrates of S. simulans (NRRL B-2628) grown
       according to published references, is an endopeptidase which cleaves the
       polyglycine cross-links of the peptidoglycan found in the cell walls of
     staphylococci. In addition, cultures that produce
     lysostaphin appear to be resistant to its activities while
       cultures grown under non-lysostaphin producing conditions are sensitive.
       Studies on the possible mechanism of antibiotic evasion of phagocytized
SUMM
     staphylococci in mastitis treatment show that
     lysostaphin had been rejected as a candidate for destroying
       phagocytized staphylococci. Craven et al., 29 Research in
       Veterinary Science 57 (1980); Craven et al., 21 Antimicrobial Agents and
       Chemotherapy 618 (1982);. . . Comp. Immun. Microbial. Infect. Dis. 447 (1982)) Craven et al., 51 Journal of Dairy Research 513 (1984). In
       these experiments lysostaphin was used in vitro as a
       pretreatment to destroy extracellular staphylococci prior to
       exposing the phagocytized staphylococci to cloxacillin,
       gentamicin or lysostaphin. Craven et al.'s results strongly
       suggest that lysostaphin would have no effect on mastitis
       since intracellular staphylococci were still viable after 20
       hours of incubation in a lysostaphin containing solution. 51
       Journal of Dairy Research at 515-516, and Table 2.
                of chronic nasal staphylococcal infections (Quickel, Jr. et
SUMM
       al., 22 Applied Microbiology 446 (1971)). In one case of a resistant
     staphylococcal infection, lysostaphin was given
       systemically (Stark et al., 291 Medical Intelligence 239 (1974)). In
       general, however, there has been great skepticism and.
          . . amount of lysostaphin, with or without surfactant, EDTA,
SUMM
       penicillin or other potentiating agents, are used to achieve elimination
       of the staphylococcal infection. Preferably such infusions
       contain between 2 to 400mg lysostaphin when no potentiating
       agents are present. In combinations containing potentiating agents, the
       required effective doses of lysostaphin can be lowered.
             . by conventional antibiotic (e.g. penicillin) therapy. In
SUMM
       addition, penicillin and other similar acting substances may also be
       useful together with lysostaphin as an agent against
     staphylococcal infection and contamination.
          . . strain 00 containing a recombinant plasmid which directs the
DETD
        synthesis of lysostaphin, as this provides for both high levels of
      lysostaphin production substantially free from
      staphylococcal immunogenic contaminants and facile
      lysostaphin purification since the lysostaphin
        accumulates directly in the growth medium. Bacillus sphaericus
        transformants containing the plasmid pBC16-1L have been found to be
       particularly suited.
                 developed either chronic or acute staphylococcal bovine
DETD
       mastitis despite prophylactic treatment. A single dose of from 2 to 400
       mg lysostaphin per milk gland will eliminate the infection and
        cure staphylococcal mastitis in most instances. Additional
        doses of lysostaphin may be indicated where the infection is
```

09/120030

persistent. Doses significantly higher than 400 mg are not recommended

```
as they can.
      Table IC demon stes the synergistic effect of
DETD
       /penicillin combinations on three strains of staphylococci.
       Depending on the doses of each, the combinations of lysostaphin
       plus penicillin can be 100 to 1000 times more effective than either
       lysostaphin or penicillin alone with all three strains.
              lysostaphin which were sufficient to eliminate the infection
DETD
       did not produce adverse side effects and indicated that intramammary
       infusions of lysostaphin are effective against
     staphylococcal mastitis. At 125 .mu.g/kg, glands were cleared of
       infection by the 6 hour post-treatment sample and remained clear
       throughout the.
                     TABLE IV
DETD
Efficacy of Intramammary Infusion of Lysostaphin Toward
Experimental STAPHYLOCOCCAL Mastitis in Guinea Pig
         Lysostaphin Dose .mu.g/kg
                           5.0 25.0 62.5 125.0
                    1.0
Number of animals
                    (1/0) (1/2) (2/2)
           (0/10)
                                      (1/1) (7/7)
cleared of
infection
       It can be seen from these examples that lysostaphin is
DETD
       effective for treatment of staphylococcal mastitis and that
       its effect is greatly enhanced when used in combination with penicillin
       or with substances such as mild. .
L13 ANSWER 4 OF 17 USPATFULL
       US 5708160 980113
PΤ
       1998:4758 USPATFULL
ΑN
       HSP-60 genomic locus and primers for species identification
ΤI
       Goh, Swee Han, Vancouver, Canada
IN
       Chow, Anthony W., West Vancouver, Canada
       Hemmingsen, Sean, Saskatoon, Canada
       The National Research Council, Ottawa, Canada (non-U.S. corporation)
PΑ
       University of British Columbia, Vancouver, Canada (non-U.S. corporation)
       US 5708160 980113
PΙ
       US 95-429121 950426 (8)
ΑI
       Utility
EXNAM Primary Examiner: Jones, W. Gary; Assistant Examiner: Fredman, Jeffrey
       Fish & Richardson, P.C.
LREP
       Number of Claims: 10
CLMN
       Exemplary Claim: 1,3
ECL
       9 Drawing Figure(s); 7 Drawing Page(s)
DRWN
LN.CNT 1165
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . (Ausubel, et al., Current Protocols in Molecular Biology, Unit
DETD
       2.4.1.-2.4.2., Greene Publishing Assoc. Inc., J. Wiley and Sons, Inc.).
       For staphylococci, lysostaphin (Sigma or recombinant
       product from Applied Microbiology, Inc., New York) was substituted for
       lysozyme in facilitating cell lysis. DNA concentration. .
L13 ANSWER 5 OF 17 USPATFULL
       US 5702895 971230
PΙ
       97:123039 USPATFULL
ΑN
       Method and kit for detecting methicillin-resistant Staphylococcus aureus
TI
       Matsunaga, Hironari, Hiroshima, Japan
IN
       Tsukumo, Kenichi, Hiroshima, Japan
```

Wakisaka, Shinji, Hiroshima, Japan Yamane, Akio, Hiroshima, Japan

```
Wakunaga Selyaku Kabushiki Kalsha, Osaka, Japan (non-o.s. colpolucion-
PΑ
      us 5702895 97 0
us 96-586274 560116 (8)
PΙ
ΑI
      JP 95-6390 950119
PRAI
DT
      Utility
EXNAM Primary Examiner: Sisson, Bradley L.
      Oblon, Spivak, McClelland, Maier & Neustadt, P.C.
LREP
      Number of Claims: 11
CLMN
      Exemplary Claim: 1
ECL
DRWN
      No Drawings
LN.CNT 747
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . a method using a protease such as proteinase K may also be
       employed. When enzymes which lyse membranes characteristic to
     Staphylococci such as lysostaphin and achromopeptidase
       are used, these enzymes enhance efficiency of extracting nucleic acids
       and elevate sensitivity of the examination.
L13 ANSWER 6 OF 17 USPATFULL
       US 5587288 961224
PΙ
       96:118502 USPATFULL
ΑN
       Regulation of exoprotein in Staphylococcus aureus
ΤI
       Cheung, Ambrose, New York, NY, United States
IN
       Fischetti, Vincent A., West Hempstead, NY, United States
       The Rockefeller University, New York, NY, United States (U.S.
PA
       corporation)
       US 5587288 961224
ΡI
       US 94-248505 940524 (8)
ΑI
       Utility
EXNAM Primary Examiner: Jones, W. Gary; Assistant Examiner: Fredman, Jeffrey
       Burns, Doane, Swecker & Mathis, L.L.P.
LREP
       Number of Claims: 3
CLMN
       Exemplary Claim: 1
ECL
       3 Drawing Figure(s); 2 Drawing Page(s)
DRWN
LN.CNT 506
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       More specifically, the suspected staphylococcal isolate may be
DETD
       incubated with lysostaphin to digest the cell and release its
       DNA which is purified by any of the procedures known to the skilled.
L13 ANSWER 7 OF 17 USPATFULL
       US 5496706 960305
PΤ
       96:18980 USPATFULL
ΑN
       Methods and materials for the detection of Staphylococcus aureus
ΤI
       Kuusela, Pentti, Helsinki, Finland
IN
       Hilden, Pekka, Helsinki, Finland
       Helsinki University Licensing, Ltd., Helsinki, Finland (non-U.S.
PΆ
       corporation)
PΤ
       US 5496706 960305
       US 93-169524 931217 (8)
AΙ
       Utility
DT
EXNAM Primary Examiner: Scheiner, Toni R.; Assistant Examiner: Parsons, Nancy
       J.
       Marshall, O'Toole, Gerstein, Murray & Borun
LREP
       Number of Claims: 6
CLMN
       Exemplary Claim: 1
ECL
       4 Drawing Figure(s); 3 Drawing Page(s)
DRWN
LN.CNT 565
       The present invention provides an approximately 230 kDa protein,
SUMM
       designated MRSA-230, which is isolated from lysostaphin
       digests of methicillin-resistant Staphylococcus aureus which
```

test negative in standard S. aureus agglutination assays. Anti-MRSA-230

```
L13 ANSWER 8 OF 17 USPATFULL
      US 5437978 950801
PΙ
      95:69206 USPATFULL
ΝA
      Detection for Staphylococcus spp.
ΤI
      Ubukata, Kimiko, Tokyo, Japan
IN
      Nakagami, Satoru, Hiroshima, Japan
      Yamane, Akio, Miyoshi, Japan
      Wakunaga Seiyaku Kabushiki Kaisha, Osaka, Japan (non-U.S. corporation)
PΑ
      US 5437978 950801
PΙ
      US 92-924458 920804 (7)
ΑI
       JP 91-195398 910805
PRAI
DT
       Utility
      Primary Examiner: Patterson, Jr., Charles L.; Assistant Examiner: Kim,
EXNAM
       Hyosuk
       Bacon & Thomas
LREP
       Number of Claims: 20
CLMN
       Exemplary Claim: 1
ECL
       2 Drawing Figure(s); 2 Drawing Page(s)
DRWN
LN.CNT 825
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
            . enzymes for hydrolyzing proteins of the bacteria. Any enzymes
DRWD
       for lysing cell walls are available which can hydrolyze peptidoglycans
       of Staphylococcus spp., and so, for example,
     lysostaphin, acromopeptidase and the like can be used. On the
       other hand, any enzymes for hydrolyzing proteins are available which
       can.
L13 ANSWER 9 OF 17 USPATFULL
       US 5342612 940830
PΙ
       94:75297 USPATFULL
ΑN
       Compositions for the treatment of mammalian diseases
ΤI
       Daley, Michael J., Yardley, PA, United States
IN
       Steber, William D., Ledgewood, NJ, United States
       Furda, Gary J., Trenton, NJ, United States
       Johnston, Paul A., Langhorne, PA, United States
       Oldham, Elizabeth R., Newtown, PA, United States
       American Cyanamid Company, Wayne, NJ, United States (U.S. corporation)
PA
       US 5342612 940830
PΙ
       US 91-812894 911220 (7)
ΑI
DT
       Utility
EXNAM Primary Examiner: Wityshyn, Michael G.; Assistant Examiner: Sayala, C.
       Morris, Michael P.
LREP
       Number of Claims: 1
CLMN
       Exemplary Claim: 1
ECL
DRWN
       No Drawings
LN.CNT 911
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . Med., 39:230 (1967) and bovine mastitis caused by S. aureus
SUMM
        (Sears et al., J. Dairy Science, 71 (Suppl. 1): 244(1988)).
     Lysostaphin, a gene product of Staphylococcus
       simulans, exerts a bacteriostatic and bactericidal effect upon S. aureus
       by enzymatically degrading the polyglycine crosslinks of the cell wall.
       Efficacy of Lysostaphin Formulated in Vehicles as an
DETD
       Intramammary Infusion Preparation Against Staphylococcus
       Aureus Mastitis
L13 ANSWER 10 OF 17 USPATFULL
       US 5011772 910430
 PΤ
```

Antibodies or antisera comprising the antibodies are useru., ......

91:34292 USPATFULL

ΑN

```
High yield protein production system
Recsei, Paul F New York, NY, United States
ΤI
IN
      Public Health Research Institute of the City of N.Y., New York, NY,
PΑ
       United States (U.S. corporation)
       US 5011772 910430
ΡI
       US 88-152635 880205 (7)
ΑI
       Utility
DT
      Primary Examiner: Martinell, James
EXNAM
       White & Case
LREP
       Number of Claims: 34
CLMN
       Exemplary Claim: 1
ECL
       7 Drawing Figure(s); 5 Drawing Page(s)
DRWN
LN.CNT 909
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Lysostaphin is a bacteriocin which lyses staphylococci
       . Plasmid pRG5 containing a 1.5 Kb cloned DNA fragment which codes for
       preprolysostaphin has been described in U.S. patent application.
L13 ANSWER 11 OF 17 USPATFULL
       US 4980163 901225
PΙ
       90:98514 USPATFULL
ΑN
       Novel bacteriocin compositions for use as enhanced broad range
ΤI
       bactericides and methods of preventing and treating microbial infection
       Blackburn, Peter, New York, NY, United States
IN
       Gusik, Sara-Ann, New York, NY, United States
       Polak, June, New York, NY, United States
       Rubino, Stephen D., New York, NY, United States
       Public Health Research Institute of the City of New York, New York, NY,
PΑ
       United States (U.S. corporation)
       US 4980163 901225
PI
       US 89-317627 890301 (7)
AΙ
       Utility
DT
EXNAM Primary Examiner: Schain, Howard E.; Assistant Examiner: Koh, Choon
       White & Case
LREP
       Number of Claims: 24
CLMN
       Exemplary Claim: 1
ECL
       No Drawings
DRWN
LN.CNT 445
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . component is present in the enhanced broad range bactericide in
       sufficient amount such that the bactericide is more effective against
     staphylococci than is lysostaphin alone and is more
       effective at treating and preventing a broad range of microbial
        infections. Methods of treating bacterial infections.
               related to the species of their origin. Lysostaphin is a
SUMM
       bacteriocin that lyses and kills practically all known species of
     Staphylococcus, but is inactive against bacteria of other
        genera. Lysostaphin, isolated from culture filtrates of
     Staphylococcus simulans (NRRL B-2628) grown according to
       published references, is an endopeptidase which cleaves the polyglycine
        cross-links of the peptidoglycan found.
        Lysostaphin is a naturally occurring bacteriocin secreted by a single
 SUMM
        known strain of S. simulans originally isolated and named
      Staphylococcus staphylolyticus by Schindler and Schuhardt. The
        production of lysostaphin by S. staphylolyticus has been
        described previously in U.S. Pat. No. 3,278,378 issued Oct. 11, 1966 and
        in Proceedings of.
        . . . effective as a bactericide towards Staphylococcus, and nisin is
 SUMM
        present in an amount sufficient to enhance the bactericidal effect of
      lysostaphin toward Staphylococci. Other compositions
        comprise lysostaphin, nisin, and a chelating agent and may
        also contain a surfactant. This composition in a carrier yields a novel
```

bactericide.

```
the bacteriocin nisin in the range of 0.1 to 300 .mu.g/mi ana
                        tericide is significantly more pactericidal towards
     the resulting tericide is significantly mor staphylococcus then lysostaphin alone. The total
SUMM
      bactericidal activity of such a novel bactericide is believed to be
       further potentiated and effective against a.
               strain 00, containing a recombinant plasmid which directs the
       synthesis of lysostaphin. This provides for production of high levels of
     lysostaphin substantially free from staphylococcal
SUMM
       immunogenic contaminants and facile lysostaphin purification
       since the lysostaphin accumulates directly in the growth
       medium. B. sphaericus transformants containing plasmids pBC16-lL or
       pROJ6649-IL have been found to be particularly.
       Nisin alone in milk has little practical bactericidal activity towards
        bactericidal towards S. aureus and can produce more than a five log
      Staphylococci. Lysostaphin alone in milk is
 DETD
        reduction in viable cells.
 L13 ANSWER 12 OF 17 USPATFULL
        US 4931390 900605
 ΡI
        90:44457 USPATFULL
         Expression of the cloned lysostaphin gene
 ΝA
         Recsei, Paul A., New York, NY, United States
         Public Health Research Institute of the City of New York, Inc., New
  ΤI
  IN
         York, NY, United States (U.S. corporation)
  PA
         Continuation-in-part of Ser. No. US 86-852407, filed on 16 Apr 1986, now
  PΙ
  _{\rm IA}
  RLI
          abandoned
  EXNAM Primary Examiner: Martinell, James
          White & Case
          Number of Claims: 23
   LREP
          Exemplary Claim: 1
   CLMN
          1 Drawing Figure(s); 1 Drawing Page(s)
   \mathtt{ECL}
   DRWN
          The present invention provides recombinant plasmids which is
   CAS INDEXING IS AVAILABLE FOR THIS PATENT.
          transformant microbial hosts express lysostaphin, a
          bacteriocin that kills most known staphylococcal species. The
           invention also provides lysostaphin, substantially free from
           non-lysostaphin contaminants. Recombinant plasmids, pRG5,
           pJP1, pDF8 and pRP1, were derived by inserting a 1.5 kilobase segment of
                 application Ser. No. 852,407, filed Apr. 16, 1986, now
           abandoned. The present invention relates to novel plasmids which in
           transformant microbial hosts express the gene for
    PARN
         lysostaphin. The invention also relates to lysostaphin so
            Lysostaphin is a bacteriocin secreted by a single known strain
            of Staphylococcus simulans orgininally isolated and named
          Staphylococcus staphylolyticus by Schindler and Schuhardt. The
     PARN
            production of lysostaphin by S. staphylolyticus has been
            described previously in U.S. Pat. No. 3,278,378 issued Oct. 11, 1966 and
            Bacteriocins are proteins secreted by bacteria that kill and sometimes
            lyse related bacteria. For example, lysostaphin lyses and
            kills practically all known staphylococcal species but is
     PARN
            inactive against bacteria of all other genera. Although its catalytic
            properties are not well characterized, lysostaphin has.
             . . . and purification of lysostaphin by known techniques, however,
             results in a product that is contaminated to some degree by other
           staphylococcal products. Immunization of animals or man with
      PARN
           lysostaphin contaminated by non-lysostaphin
```

```
immunogenic material from staphylococci might result in an undesirable, potentially adverse, immunological response.

25,000 daltons. It is heat labile, no lalyzable and has an
       isoelectric point of about pH 11. Futhermore, the capacity of
    lysostaphin to lyse viable and heat-killed staphylococci
       and staphylococcal cell walls is destroyed by treatment with
PARN
       In accordance with the present invention, recombinant plasmids are
       described which in transformant microbial hosts will express a
       gene encoding lysostaphin. The recombiant plasmids were
       derived by inserting an identified DNA sequence which codes for
PARN
        lysostaphin into suitable cloning vectors.
       The lysostaphin expressed as a result of transformation of
      microbial hosts by the above-mentioned plasmids and other
        plasmids containing the lysostaphin gene is substantially free
 PARN
        of non-lysostaphin contaminants, especially immunogenic
        The invention also provides for preprolysostaphin, prolysostaphin and
      staphylococcal contaminants.
      lysostaphin, which is substantially free of non-
      lysostaphin immunogenic staphylococcal contaminants.
        The invention further encompasses those portions of the 1.5 kbp DNA
 PARN
         fragment which code for the lysostaphin signal peptide, . .
            immunologic cross-reactivity with lysostaphin-specific
         antibodies, catalytic activity. Lysostaphin produced by transformant
         microorganisms according to this invention is substantially free of non-
  DETD
       lysostaphin contaminants, in particular immunogenic
         Approximately 1% of the B. subtilis cells transformed with the ligated
        staphylococcal contaminants.
          DNA produced lysostaphin as indicated by the lysis of
        staphylococcal cells surrounding the B. subtilis colonies. pJP1
          was obtained by restreaking one of the B. subtilis transformants on
          of B. sphaericus 00/pJP1 transformants is substantially free of
          lysostaphin indicator. .
          non-lysostaphin contaminants. Of special significance is that the B.
          sphaericus 00/pJP1 lysostaphin is substantially free of
   DETD
           immunogenic staphylococcal contaminants.
           1. Recombinant plasmids containing a DNA sequence which codes for
           What is claimed is:
         lysostaphin and which in transformant microbial hosts
    CLM
           will express a gene encoding lysostaphin from S. simulans
            (NRRLB-2628).
    L13 ANSWER 13 OF 17 USPATFULL
            US 4902616 900220
            Process for the preparation of capsular polysaccharides of
     PΙ
            staphylococci, the polysaccharides obtained, uses of these
     ΑN
            polysaccharides and strains for carrying out of the process
     ΤI
            Fournier, Jean-Michel, Paris, France
            Bouvet, Anne, Paris, France
     IN
             Institut Pasteur, Paris, France (non-U.S. corporation)
            Boutonnier, Alain, Paris, France
             US 4902616 900220
      PΑ
             US 88-227137 880802 (7)
      PΙ
             Primary Examiner: Griffin, Ronald W.; Assistant Examiner: Webber, Pamela
      ΑI
      PRAI
      EXNAM
              Burns, Doane, Swecker & Mathis
              Number of Claims: 15
       LREP
              Exemplary Claim: 1
       CLMN
       ECL
              No Drawings
```

DRWN

```
specific enzyme for staphylococcus such as lysostaphin
    ANSWER 14 OF 17 USPATFULL
      US 4810567 890307
AN
       89:17168 USPATFULL
      Antimicrobial fabrics utilizing graft copolymers
TI
       Calcaterra, Lidia T., Des Plaines, IL, United States
IN
       DeFilippi, Louis J., Mt. Prospect, IL, United States
       Childs, Michael E., Medford, NJ, United States
       Latos, Edwin J., Chicago, IL, United States
       UOP, Des Plaines, IL, United States (U.S. corporation)
       US 4810567 890307
      US 87-94767 870910 (7)
ΑI
       Continuation-in-part of Ser. No. US 85-768090, filed on 21 Aug 1985, now
       abandoned
      Utility
EXNAM Primary Examiner: Bell, James J.
       McBride, Thomas K.; Snyder, Eugene I.
CLMN
       Number of Claims: 19
       Exemplary Claim: 14
ECL
      No Drawings
DRWN
LN.CNT 945
       Fabrics containing bound lysostaphin were tested for
     antimicrobial properties on V-J agar media. In this test five
       1-cm.sup.2 pieces of fabric were placed on V-J agar to which.
L13 ANSWER 15 OF 17 USPATFULL
       US 4801449 890131
PΙ
       89:7415 USPATFULL
ΑN
       Method for treatment of Kaposi's sarcoma
       Balint, Jr., Joseph P., Seattle, WA, United States
IN
       Jones, Frank R., Edmonds, WA, United States
       IMRE Corporation, Seattle, WA, United States (U.S. corporation)
PΑ
       US 4801449 890131
PΙ
       US 86-948268 861231 (6)
ΑI
       Continuation-in-part of Ser. No. US 85-690781, filed on 11 Jan 1985, now
RLI
       patented, Pat. No. US 4681870
       Utility
DT
EXNAM Primary Examiner: Kight, John; Assistant Examiner: Nutter, Nathan M.
       Townsend & Townsend
LREP
       Number of Claims: 10
CLMN
       Exemplary Claim: 1
ECL
       2 Drawing Figure(s); 2 Drawing Page(s)
DRWN
LN.CNT 544
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . to silica prepared as described above, and returned to the
DETD
       patient. The protein A was isolated from pure cultures of
     Staphylococcus aureus Cowan I employing lysostaphin
       digestion. Protein A purity was determined by polyacrylamide gel
       electrophoresis, and IgG binding capacity was determined. The protein A
       was. .
L13 ANSWER 16 OF 17 USPATFULL
       US 4783484 881108
PΙ
       88:72431 USPATFULL
       Particulate composition and use thereof as antimicrobial agent
TI
       Violante, Michael R., Rochester, NY, United States
IN
       Steigbigel, Roy T., Miller Pl., NY, United States
```

after bacterial lysis, brought about by autoclaning or by the use of a

LN.CNT 369

PA

CAS INDEXING IS AVAIL FOR THIS PATENT.

University of Rochester, Rochester, NY, United States (U.S. corporation)

```
US 4783484 881
PΙ
                         05 (6)
       US 84-658153 8
AΤ
       Utility
DΨ
      Primary Examiner: Brown, J. R.; Assistant Examiner: Rollins, Jr., John
EXNAM
      Kenyon & Kenyon
LREP
      Number of Claims: 29
CLMN
ECL
      Exemplary Claim: 1
DRWN
      No Drawings
LN.CNT 899
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . . leukocytes were centrifuged at 200 g for 10 minutes, decanted,
DETD
       and the pellet resuspended in PBS containing 10 U/ml of
     lysostaphin for lysis of the remaining extracellular
     Staphylococci. After 10 minutes in 37.degree. C. water bath,
       tubes were centrifuged 10 minutes at 200 g, decanted and the pellet. .
L13 ANSWER 17 OF 17 USPATFULL
      US 4681870 870721
       87:52185 USPATFULL
ΑN
       Protein A-silica immunoadsorbent and process for its production
ТT
       Balint, Jr., Joseph P., Seattle, WA, United States Hargreaves, Richard E., Seattle, WA, United States
ΙN
       IMRE Corporation, Seattle, WA, United States (U.S. corporation)
PA
       US 4681870 870721
ΡI
       US 85-690781 850111 (6)
       Utility
EXNAM Primary Examiner: Garvin, Patrick P.
       Townsend & Townsend
LREP
       Number of Claims: 26
CLMN
ECL
       Exemplary Claim: 1
       2 Drawing Figure(s); 2 Drawing Page(s)
DRWN
LN.CNT 616
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       . . to silica prepared as described above, and returned to the
DETD
       patient. The protein A was isolated from pure cultures of
     Staphylococcus aureus Cowan I employing lysostaphin
       digestion. Protein A purity was determined by polyacrylamide gel
       electrophoresis, and IgG binding capacity was determined. The protein A
       was.
=> d 13 pi,bib, kwic 1-17
L23 ANSWER 13 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
                                                        DUPLICATE 8
     1997:199416 BIOSIS
NA
     PREV199799498619
DN
     Studies on prolysostaphin processing and characterization of the
     lysostaphin immunity factor (Lif) of Staphylococcus
     simulans biovar staphylolyticus.
     Thumm, Guether; Goetz, Friedrich (1)
     (1) Mikrobielle Genet., Univ. Tuebingen, Waldhaeuser Stasse 708, D-72076
     Tuebingen Germany
     Molecular Microbiology, (1997) Vol. 23, No. 6, pp. 1251-1265.
     ISSN: 0950-382X.
     Article
DT
LA
     English
     Studies on prolysostaphin processing and characterization of the
     lysostaphin immunity factor (Lif) of Staphylococcus
     simulans biovar staphylolyticus.
     Lysostaphin is an extracellular glycylglycine endopeptidase
AΒ
```

produced by Staphylococcus simulans biovar staphylolyticus

```
ATCC1362 that lys staphylococcal cells by hydrolysing the polygraminterpeptide brid of the peptidoglycan. Renewed analysis of. . .
L23 ANSWER 1 OF 280 CAPLUS COPYRIGHT 1999 ACS
     PATENT NO. KIND DATE APPLICATION NO. DATE
     US 5760026 A 19980602 US 94-303551 19940909
US 5858962 A 19990112 US 93-168687 19931216
PΙ
     1998:392144 CAPLUS
AN
     129:23422
DN
     Method using lysostaphin for treating mastitis and other
     staphylococcal infections
     Blackburn, Peter; Polak, June
IN
     Ambi Inc., USA
PA
     U.S., 10 pp. Cont. of U. S. Ser. No. 935121, abandoned.
     CODEN: USXXAM
DT
     Patent
LΑ
     English
FAN.CNT 1
                                             APPLICATION NO. DATE
     PATENT NO. KIND DATE
     -----
PI US 5760026 A 19980602 US 94-303551 19940909
US 5858962 A 19990112 US 93-168687 19931216

PRAI US 87-48412 19870511
US 88-188183 19880428
US 90-535286 19900608
US 92-935121 19920820
US 89-440092 19891122
                                              _____
     Method using lysostaphin for treating mastitis and other
     staphylococcal infections
     Lysostaphin is used to eliminate and cure staphylococcal
\mathbf{A}\mathbf{B}
     infections including the cure of mastitis by intramammary infusion.
     Administration of 2-400 mg of lysostaphin to an infected bovine
     mammary gland eliminates staphylococci, and the reoccurrence
     common with antibiotic therapy is not obsd. Teat-dips contg. lysostaphin,
     mutanolysin and lysozyme can be used as.
     lysostaphin mastitis staphylococcal infection; synergy
     lysostaphin surfactant mastitis staphylococcal
     infection; penicillin lysostaphin synergy mastitis
     staphylococcal infection
     Plasmids
IT
         (PBC16-1L; recombinant lysostaphin for treating mastitis and
         other staphylococcal infections)
     Escherichia coli
     Klebsiella pneumoniae
     Mastitis
     Milk
      Staphylococcus aureus
      Staphylococcus epidermidis
     Streptococcus agalactiae
      Streptococcus uberis
         (lysostaphin for treating mastitis and other
       staphylococcal infections)
      Bacillus sphaericus
IT
         (lysostaphin prodn. in; recombinant lysostaphin for
         treating mastitis and other staphylococcal infections)
ΙT
      Antibacterial agents
      Chelating agents
      Surfactants
      Synergistic drug interactions
```

(lysostaphin, and synergistic combinations, for treating

mastitis and other staphylococcal infections)

ΙT

Genes

```
RL: BPR (Biologic process); BIOL (Biological study); PROC (FIOCESS) — (lysostaphin; ombinant lysostaphin for treating
       mastitis and other staphylococcal infections)
    9011-93-2P, Lysostaphin
ΙT
    RL: BAC (Biological activity or effector, except adverse); BPN
     (Biosynthetic preparation); THU (Therapeutic use); BIOL (Biological
     study); PREP (Preparation); USES (Uses)
        (lysostaphin for treating mastitis and other
     staphylococcal infections)
     55466-22-3, Mutanolysin
ΙT
    RL: BAC (Biological activity or effector, except adverse); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (lysostaphin for treating mastitis and other
     staphylococcal infections)
     9001-\overline{63}-2, Lysozyme
IT
     RL: BAC (Biological activity or effector, except adverse); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (lysostaphin, and combinations, for treating mastitis and
        other staphylococcal infections)
     61-32-5, Methicillin 1405-87-4, Bacitracin 1406-05-9, Penicillin 1406-11-7, Polymyxin 9002-93-1, Triton X-100 11111-12-9, Cephalosporin
IT
     RL: BAC (Biological activity or effector, except adverse); THU
     (Therapeutic use); BIOL (Biological study); USES (Uses)
        (lysostaphin, and synergistic combinations, for treating
        mastitis and other staphylococcal infections)
     9073-60-3, Penicillinase
IT
     RL: BSU (Biological study, unclassified); BIOL (Biological study)
        (penicillinase-pos. Staphylococcus aureus;
      lysostaphin for treating mastitis and other
      staphylococcal infections)
L23 ANSWER 2 OF 280 CAPLUS COPYRIGHT 1999 ACS
    PATENT NO. KIND DATE APPLICATION NO. DATE
     ______
<---->
     Preparation of the luciferase-lysostaphin fusion protein for
     detection of Staphylococcus aureus by bioluminescence analysis
     Tatsumi, Hiroki; Fukuda, Masaru; Nagahara, Ayumu
IN
     Kikkoman Corp., Japan
PA
     Jpn. Kokai Tokkyo Koho, 10 pp.
     CODEN: JKXXAF
DT
    Patent
LΑ
    Japanese
FAN.CNT 1
     PATENT NO. KIND DATE APPLICATION NO. DATE
                                          -----
     JP 10150991 A2 19980609 JP 96-328042 19961125
     Preparation of the luciferase-lysostaphin fusion protein for
     detection of Staphylococcus aureus by bioluminescence analysis
     Disclosed are a fusion protein comprised of luciferase[217-Leu] of Luciola
     lateralis and the C-terminal fragment of lysostaphin of
     Staphylococcus simulans (NRRL B-2628) by expression of its
     encoding chimeric gene in Escherichia coli and use of the fusion protein
     luciferase lysostaphin fusion protein Escherichia prepn;
ST
     Staphylococcus detection luciferase lysostaphin fusion
     Genes (microbial)
IT
     RL: BPN (Biosynthetic preparation); BUU (Biological use, unclassified);
     PRP (Properties); BIOL (Biological study); PREP (Preparation); USES (Uses)
         (chimeric; prepn. of luciferase-lysostaphin fusion protein
```

for detection of Staphylococcus aureus by bioluminescence